

GenCore version 4.5
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TM nucleic - nucleic search, using sw model

Run on: February 18, 2001, 08:51:31 ; Search time 128.56 Seconds

(without alignments)
8643.515 Million cell updates/sec

Title: US-09-434-382-3

Perfect score: 2958

Sequence: 1 cggggcgtagtgaccggc.....aataaagattgagtttgcac 2958

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 480022 seqs, 187831343 residues

Word size : 0

Total number of hits satisfying chosen parameters: 960044

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 100 summaries

Database : N_Geneseq_36.*

1: /cgnl_8/gcgdata/geneseq/geneseq/NA1980.DAT.*
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3: /cgnl_8/gcgdata/geneseq/geneseq/NA1982.DAT.*
4: /cgnl_8/gcgdata/geneseq/geneseq/NA1983.DAT.*
5: /cgnl_8/gcgdata/geneseq/geneseq/NA1984.DAT.*
6: /cgnl_8/gcgdata/geneseq/geneseq/NA1985.DAT.*
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20: /cgnl_8/gcgdata/geneseq/geneseq/NA1999.DAT.*
21: /cgnl_8/gcgdata/geneseq/geneseq/NA2000.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	237	8.0	238	21	280231
2	53	1.8	72	16	T25953
3	39	1.3	68	19	X11554
4	19	0.6	526	20	V88751
5	19	0.6	7642	20	V82020
6	18	0.6	45	20	Z33905
7	18	0.6	177	20	X18065
8	18	0.6	531	18	T67773
9	18	0.6	531	18	T77453
10	18	0.6	726	21	Z80749
11	18	0.6	1040	21	Z48812
12	18	0.6	1593	18	T67992
					Human colon cancer
					Human gene signatu
					Human biallelic po
					EST clone HK189.
					Moraxella catarrha
					Human PRO274 hybri
					Coding sequence fo
					H. pylori cytoplas
					H. pylori cytoplas
					Human colon cancer
					Soybean inositol 1
					H. pylori cytoplas

c 13	18	0.6	1626	20	X08683
c 14	18	0.6	2186	19	V17351
c 15	18	0.6	2450	20	X20537
c 16	18	0.6	2605	19	V04699
c 17	18	0.6	2945	20	X23895
c 18	18	0.6	4120	20	Z09473
c 19	18	0.6	6139	19	V70354
c 20	18	0.6	49999	20	Z23903
c 21	17	0.6	21	15	Q54684
c 22	17	0.6	256	21	A32079
c 23	17	0.6	300	21	A32050
c 24	17	0.6	300	21	A01382
c 25	17	0.6	304	17	T07238
c 26	17	0.6	304	17	Z42643
c 27	17	0.6	332	20	X24993
c 28	17	0.6	347	16	T24629
c 29	17	0.6	369	20	X41005
c 30	17	0.6	422	20	X24994
c 31	17	0.6	426	21	A32069
c 32	17	0.6	431	21	A32052
c 33	17	0.6	559	19	V29360
c 34	17	0.6	590	20	X24995
c 35	17	0.6	605	20	V87198
c 36	17	0.6	791	20	X24979
c 37	17	0.6	855	21	Z91823
c 38	17	0.6	1022	20	X40199
c 39	17	0.6	1084	15	Q78866
c 40	17	0.6	1084	20	Z08442
c 41	17	0.6	1094	15	Q78865
c 42	17	0.6	1094	20	Z08441
c 43	17	0.6	1140	20	X27353
c 44	17	0.6	1173	13	Q29634
c 45	17	0.6	1173	13	Q29635
c 46	17	0.6	1173	14	Q43895
c 47	17	0.6	1209	21	Z46089
c 48	17	0.6	1338	20	X87593
c 49	17	0.6	1341	20	X87591
c 50	17	0.6	1345	12	Q14450
c 51	17	0.6	1557	17	T27644
c 52	17	0.6	1591	19	V98595
c 53	17	0.6	1592	19	V42963
c 54	17	0.6	1632	19	V68059
c 55	17	0.6	1664	20	Z25023
c 56	17	0.6	1668	20	V72116
c 57	17	0.6	1891	20	V69719
c 58	17	0.6	1702	20	Z52876
c 59	17	0.6	1724	19	V38385
c 60	17	0.6	1816	20	X85940
c 61	17	0.6	1818	20	X01577
c 62	17	0.6	1906	21	A15550
c 63	17	0.6	2160	16	T09084
c 64	17	0.6	2284	21	Z50582
c 65	17	0.6	2418	20	X84103
c 66	17	0.6	2419	13	Q32351
c 67	17	0.6	2419	15	Q72476
c 68	17	0.6	2419	16	T05086
c 69	17	0.6	2419	20	X84112
c 70	17	0.6	2420	15	Q72472
c 71	17	0.6	2420	16	Q85435
c 72	17	0.6	2503	19	V59595
c 73	17	0.6	2646	20	Z42096
c 74	17	0.6	2711	19	V68056
c 75	17	0.6	2971	9	N81166
c 76	17	0.6	3000	12	Q13115
c 77	17	0.6	3156	19	V18471
c 78	17	0.6	3156	20	Z27969
c 79	17	0.6	3457	14	Q48468
c 80	17	0.6	3796	18	T93439
c 81	17	0.6	4216	8	N70558
c 82	17	0.6	4308	18	T45351
c 83	17	0.6	4308	21	Z51806
c 84	17	0.6	4488	14	Q51426
c 85	17	0.6	4567	20	V33945

Novel nucleotide s
Coding sequence fo
Polynucleotide seq
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Human PRO274 nucle
Human RNA helicase
Coding strand of n
Human LOBO homolog
C kappa exon prime
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Plant microsatelli
Immunogen DNA from
Human 5' EST isola
Murine Bcl-2 inter
Human gene signatu
Human secreted pro
Murine Bcl-2 inter
Plant microsatelli
Plant microsatelli
Calcium ion channe
Murine Bcl-2 inter
EST clone BN180.
Streptococcus pneu
MAGE-4 encoding ge
H6/MAGE-1 expressi
H6/MAGE-1 expressi
H6/MAGE-1 expressi
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Hepatitis C virus
Hepatitis C virus
CDNA encoding a fo
CLYTA-MAGE-1-His f
Lipoprotein D-MAGE
16S RNA from ATCC
CDNA encoding prot
DNA encoding GTP-b
Streptococcus pneu
Neurodegenerative
Murine D6 encoding
Mouse FAST-1 codin
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Human prostate tum
Beta(1 -> 4)-N-ace
Antigen E gene. H
Tumour rejection a
M22-MEL antigen E
Antigen E coding s
Tumour rejection a
Human melanoma ant
Human secreted pro
Human endometrium
Neurodegenerative
fdhf gene. synthe
Encodes partial mu
T-cell surface ant
Human CD97 protein
Nitrate reductase
Xenopus frog prote
Sequence of nitrog
Human colon carcin
Full length expand
Human FACC cDNA cl
Fanconi anaemia CO

86 17 0.6 5674 13 Q32352
 87 17 0.6 5674 15 Q74477
 88 17 0.6 5674 20 X84113
 89 17 0.6 5720 17 T42117
 90 17 0.6 5720 21 Z51508
 91 17 0.6 5724 16 Q98902
 92 17 0.6 6464 14 Q48772
 93 17 0.6 10240 19 V52165
 94 17 0.6 10461 20 X20553
 95 17 0.6 10813 18 V74675
 96 17 0.6 11236 15 Q70447
 97 17 0.6 21170 20 X20535
 98 17 0.6 29604 18 X83005
 99 17 0.6 34094 20 Z30163
 100 17 0.6 38734 20 Z32020

ALIGNMENTS

RESULT 1
 30231
 280231 standard; cDNA; 238 BP.
 280231;
 07-APR-2000 (first entry)
 Human colon cancer cell line SW480 cDNA clone SEQ ID NO:315.
 Human; gene expression product; diagnosis; tumour; colon cancer;
 colorectal adenocarcinoma; cell line SW480; cell proliferation;
 cytostatic; sarcoma; breast cancer; neoplasia; dysplasia;
 hyperplasia; ds.
 Homo sapiens.
 WO9964576-A2.
 16-DEC-1999.
 09-JUN-1999; 99WO-IB01062.
 10-JUN-1998; 98US-0088801.
 (FARB) BAYER CORP.
 Endege WO, Steinmann KE, Astle JH, Burgess CC, Bushnell SE;
 Carroll E, Catino TJ, Derti A, Ford DM, Lewis ME, Monahan JE;
 Schlegel R;
 WPI; 2000-087220/07.
 Novel nucleic acids, used to develop products for the diagnosis and
 treatment of disorders involving unwanted cell proliferation,
 particularly cancers, especially colon cancer
 Claim 15; Page 258; 469pp; English.
 79917 to 280766 represent double stranded cDNA clones isolated from the
 human colorectal adenocarcinoma (colon cancer) cell line SW480. The
 cDNA clones can be used to generate antisense oligonucleotides which
 can be used for antisense therapy. Methods and products from the present
 invention can be used for identifying and/or classifying cancerous cells
 present in a human tumour, particularly in solid tumours, e.g. carcinomas
 and sarcomas, e.g. breast or colon cancers. The cDNA clones can be used
 for developing agents for the diagnosis and treatment of disorders
 involving unwanted cell proliferation, such as neoplasia, dysplasia or
 hyperplasia.
 Sequence 238 BP; 55 A; 57 C; 69 G; 57 T; 0 other;

Query Match 8.0%; Score 237; DB 21; Length 238;
 Best Local Similarity 100.0%; Pred. No. 4.1e-106; Indels 0; Gaps 0;
 Matches 237; Conservative 0; Mismatches 0;
 QY 226 acctgcaggtggtggcagcggttagcgggactcggcgctcgcgcgtctacgtcttctccg 285
 Db 1 acctgcaggtggtggcagcggttagcgggactcggcgctcgcgcgtctacgtcttctccg 60
 QY 286 agttcaacgggtatctcttcaactgtggagaaggcgttcagagactcagagagacaca 345
 Db 61 agttcaacgggtatctcttcaactgtggagaaggcgttcagagactcagagagacaca 120
 QY 346 agttaagggtgctgcctggacacatatctcctgacacaaatgcactgtctcaatgttg 405
 Db 121 agttaagggtgctgcctggacacatatctcctgacacaaatgcactgtctcaatgttg 180
 QY 406 ggggcttaagtgaatgattcttactttaaggaacccggcttccaaagtgtgtac 462
 Db 181 ggggcttaagtgaatgattcttactttaaggaacccggcttccaaagtgtgtac 237
 RESULT 2
 T25953
 ID T25953 standard; cDNA to mRNA; 72 BP.
 XX
 AC T25953;
 XX
 DT 28-OCT-1996 (first entry)
 XX
 DE Human gene signature HUMGS08188.
 XX
 KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;
 human; cloning; mapping; non-biased library; diagnosis; detection;
 cell typing; abnormal cell function; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO9514772-A1.
 XX
 PD 01-JUN-1995.
 XX
 PF 11-NOV-1994; 94WO-JF01916.
 XX
 PR 12-NOV-1993; 93JP-0355504.
 XX
 PA (MATS/) MATSUBARA K.
 PA (OKUBO/) OKUBO K.
 XX
 PI Matsubara K, Okubo K;
 XX
 DR WPI; 1995-206931/27.
 XX
 PS Identifying gene signatures in 3'-directed human cDNA library - e.g.
 for diagnosis of abnormal cell function, by preparing cDNA that
 reflects relative abundance of corresp. mRNA in specific human
 tissues
 XX
 PS Claim 1; Page 1967; 2245pp; Japanese.
 XX
 CC A single-stranded DNA (or its complementary strand or the corresp.
 double-stranded DNA) which comprises one of the 7837 "GS" sequences
 given in T19001-T26637 and which is able to hybridise to part of
 human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
 sequences were obtained from 3'-directed cDNA libraries prepared
 from various human tissues; synthesis of cDNA was initiated from the
 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
 untranslated sequence is unique to a particular mRNA species, almost
 all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
 is constructed so as to reflect accurately the relative abundance of
 different mRNAs in the particular tissue from which it was derived.
 CC The appearance frequency of a given GS in a cDNA library can be
 determined (esp. using primers and probes derived from the GS
 sequences) as a means of diagnosing abnormal cell function or for

CC recognising different cell types.

Q Sequence 72 BP; 25 A; 10 C; 16 G; 20 T; 1 other;

Query Match 1.8%; Score 53; DB 16; Length 72;
Best Local Similarity 100.0%; Pred. No. 4.4e-16;
Matches 53; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2906 tccgagacttaacgaaatagatttcagctgcaataaagattgattgcaaa 2958
|||||
b 19 tccgagacttaacgaaatagatttcagctgcaataaagattgattgcaaa 71

RESULT 3

X11554/c
D X11554 standard; DNA; 68 BP.

XX X11554;

TI 30-MAR-1999 (first entry)

Human biallelic polymorphic DNA fragment ESTC169.

Polymorphism: biallelic; human; forensic; paternity testing; disease;
detection; phenotypic typing; characteristic; infection; hereditary;
autoimmune disease; cancer; inflammation; drug; therapy; medication;
treatment; marker; ss.

XX Homo sapiens.

DS WO9820165-A2.

FN 14-MAY-1998.

XX 05-NOV-1997; 97WO-US203313.

XX 06-NOV-1996; 96US-0030455.

XX (WHED) WHITEHEAD INST BIOMEDICAL RES.

XX Hudson T, Lander ES, Wang D;

XX WPI; 1998-286974/25.

XX New isolated nucleic acid segments from the human genome - used for
determining polymorphic forms for use in e.g. forensics, paternity
testing or phenotypic typing for disease

XX Claim 1; Page 172; 310pp; English.

XX X10269-X12937 are human DNA fragments which contain biallelic polymorphic
markers which have been isolated using the primers represented in
X09121-X10268. The base occupying the polymorphic site is indicated by
the appropriate IUPAC-IUB ambiguity code. These fragments can be used in
methods for determining polymorphic forms in an individual for use in
e.g. forensics, paternity testing or for phenotypic typing for diseases
such as agammaglobulinemia, diabetes insipidus, Lesch-Nyhan syndrome,
muscular dystrophy, Wiskott-Aldrich syndrome, Fabry's disease, familial
hypercholesterolemia, polycystic kidney disease, hereditary
spherocytosis, von Willebrand's disease, tuberous sclerosis, hereditary
haemorrhagic telangiectasia, familial colonic polyposis, Ehlers-Danlos
syndrome, osteogenesis imperfecta, acute intermittent porphyria,
autoimmune diseases, inflammation, cancer, diseases of the nervous
system, infection by pathogenic microorganisms, and characteristics such
as longevity, appearance (e.g. baldness, obesity), strength, speed,
endurance, fertility, and susceptibility or receptivity to particular
drugs or therapeutic treatments. The isolated polymorphic nucleic acid
segments can also be used to produce medicaments for the treatment or
prophylaxis of such diseases.

XX Sequence 68 BP; 12 A; 13 C; 21 G; 21 T; 1 other;

Query Match 1.3%; Score 39; DB 19; Length 68;
Best Local Similarity 100.0%; Pred. No. 3.1e-09;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2812 tggaaacagacgagcgccctttctcttaataccagcaaa 2850
|||||
Db 62 TGGAAACAGACGGCGGCACCTTCTCTAATCCAGCAAA 24

RESULT 4

V88751/c
ID V88751 standard; cDNA; 526 BP.

XX V88751;

XX 12-FEB-1999 (first entry)

DE EST clone HK189.

XX Expressed sequence tag; secreted protein; haematopoiesis regulator;
tissue growth; activin; inhibitor; tumour invasion suppressor; EST; human;
chemotaxis; chemokinesis; haemostasis; gene therapy; thrombolysis;
receptor; ligand; anti-inflammatory; tumour inhibitor; ds.

XX Homo sapiens.

XX WO9845437-A2.

XX 15-OCT-1998.

XX 10-APR-1998; 98WO-US06956.

XX 10-APR-1997; 97US-0837312.

XX (GEMY) GENETICS INST INC.

XX Agostino MJ, Jacobs K, Lavallie ER, McCoy JM, Merberg D;
Racie LA, Spaulding V, Treacy M;

XX WPI; 1999-070078/06.

XX New polynucleotides encoding human secreted proteins - derived from
e.g. human blood, kidney, foetal lung, placenta, testes, brain,
ovary, pituitary, retina and colon cDNA libraries

XX Claim 1; Page 500; 641pp; English.

XX The present sequence represents an expressed sequence tag (EST), and is
a polynucleotide of the invention. The polynucleotides of the invention
are all secreted EST sequences isolated from a variety of human tissue
sources. The EST sequences and proteins encoded by them are predicted to
have useful biological activities which would make them suitable for
treating, preventing or ameliorating medical conditions in humans and
animals, although no supporting data is given. Suggested activities
include nutritional activity, immune stimulating or suppressing activity,
haematopoiesis regulating activity, tissue growth activity,
activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
and thrombolytic activity, receptor/ligand activity, anti-inflammatory
activity, cadherin/tumour invasion suppressor activity, tumour inhibition
activity. The EST sequences are also stated to be useful for gene
therapy.

XX Sequence 526 BP; 127 A; 138 C; 136 G; 125 T; 0 other;

Query Match 0.6%; Score 19; DB 20; Length 526;

Best Local Similarity 100.0%; Pred. No. 18;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 742 gaagagggtcaggactc 760

Db 272 GAAGAGGGGTCAGGACTC 254

WPI: 1999-009568/01.
 New proteins that bind specifically to receptors in the gastro-intestinal tract and related nucleic acid - chimaeras and antibodies, used to deliver therapeutic or diagnostic agents to, or through, the gastrointestinal tract, e.g. insulin or leuprolide
 Claim 49; Page 56; 294pp; English.
 This sequence encodes a peptide that specifically binds to the human sucrose-isomaltase complex. The invention relates to purified proteins (I) that bind specifically to at least one of the gastro-intestinal (GI) tract receptors human intestinal peptide-associated transporter (HPII), hPEP1, D2H and human sucrose-isomaltase complex (hSI). (I) provide active transport of therapeutic agents through human and animal GI tissue (into the blood) for in vivo delivery, particularly for treatment or prevention of hypertension, diabetes, osteoporosis, haemophilia, anaemia, cancer, migraine, or angina pectoris. Specifically they are used to deliver insulin or leuprolide, but many other suitable therapeutic agents are disclosed, including genes or inhibitory nucleic acid, imaging agents and antigens. (I) may also provide targeting to the GI tract. Other uses of (I) are: (i) to determine the level of specified receptors in a sample (in a binding assay); and (ii) to screen for molecules that bind (I). Immunogenic analogues or derivatives of (I) are used to raise antibodies and in immunoassays. The antibodies of (I) are used to locate, detect and measure (I), e.g. for imaging, monitoring treatment, tissue analysis etc., also for peptide purification and immobilisation.
 Sequence 177 BP; 31 A; 51 C; 53 G; 42 T; 0 other;

Query Match 0.6%; Score 18; DB 20; Length 177;
 Best Local Similarity 100.0%; Pred. No. 58;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

17 1816 agcagtagcaccacacagc 1833
 |||||
 90 AGCAGTACCACACCACT 73

RESULT 8
 T5773/c
 T67773 standard; DNA; 531 BP.
 T67773;

29-JUL-1997 (first entry)
 H. pylori cytoplasmic protein ORF 24824087.aa.
 Vaccine; prevention; treatment; infection; identification; binding compound; bacterium; life cycle; activator; bacteria; inhibitor; duodenal ulcer disease; chronic gastritis; diagnosis; cytoplasmic; ds.

Helicobacter pylori.
 Key Location/Qualifiers
 CDS 1..531
 /*tag= a
 /transl_except= (pos: 460..462, aa: xaa)
 /transl_except= (pos: 520..522, aa: xaa)
 /transl_except= (pos: 526..528, aa: xaa)
 /note= "Xaa = Unknown"

WO9640893-A1.
 19-DEC-1996.
 06-JUN-1996; 96WO-US09122.
 01-APR-1996; 96US-0630405.

PR 07-JUN-1995; 95US-0487032.
 XX (ASTR) ASTRA AB.
 XX Berglinth OT, Smith D, Mellgaard BL;
 PI WPI: 1997-052306/05.
 XX P-PSDB; W20335.
 DR
 XX
 PT Helicobacter pylori nucleic acid sequences and related
 PT polypeptide(s) - useful for vaccines to treat or prevent H. pylori
 PT infection, and to detect Helicobacter
 XX
 PS Claim 9; Page 1; 1481pp; English.
 XX
 CC The present sequence encodes a Helicobacter pylori cytoplasmic
 CC protein.
 CC The protein may be used in a vaccine to prevent or treat H. pylori
 CC infection or to identify H. pylori polypeptide binding compounds,
 CC useful as potential H. pylori life cycle activators or inhibitors.
 CC The genomic sequence of H. pylori (ATCC 55679) was determined from
 CC overlapping contigs generated by mechanically shearing the
 CC bacterial DNA. The sequences were analysed for ORF of at least 180
 CC nucleotides, and the predicted coding regions defined by computer
 CC evaluation. To identify likely H. pylori antigens for vaccine
 CC development, the amino acid sequences predicted from various ORF
 CC were analysed for significant homology to other known or exported
 CC membrane proteins. Having identified and determined the sequences
 CC of interest, particular regions can be isolated from H. pylori by
 CC PCR amplification for recombinant polypeptide production, e.g. in
 CC E. coli hosts.
 CC Note: This DNA sequence is not reproduced in the specification and
 CC has been derived from the related specification, WO9719098.
 XX
 SQ Sequence 531 BP; 175 A; 88 C; 114 G; 151 T; 3 other;

Query Match 0.6%; Score 18; DB 18; Length 531;
 Best Local Similarity 100.0%; Pred. No. 57;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1760 aaagccgcttcacccctt 1777
 |||||
 Db 311 AAAGCGCTTCACCCCTT 294

RESULT 9
 T77453/c
 ID T77453 standard; DNA; 531 BP.
 XX
 AC T77453;
 XX
 DT 11-AUG-1997 (first entry)
 XX
 DE H. pylori cytoplasmic protein ORF 24824087.aa.

Chronic gastritis; duodenal ulcer disease; activator;
 inhibitor; bacterial life cycle; vaccine; immunisation; detection;
 antisense; inhibition; cytoplasmic; Na+/H+ antiporter;
 Escherichia coli; ds.

Helicobacter pylori.
 Key Location/Qualifiers
 CDS 1..531
 /*tag= a
 /transl_except= (pos: 460..462, aa: xaa)
 /transl_except= (pos: 520..522, aa: xaa)
 /transl_except= (pos: 526..528, aa: xaa)
 /note= "Xaa = Unknown"

WO9719098-A1.
 XX

1X WPI; 1999-288272/24.
 2 P-PSDB; W85718.
 3
 4 New polynucleotides encoding secreted human proteins
 5
 6 Claim 14; Page 101-102; 136pp; English.
 7
 8 The new human secreted proteins are encoded by polynucleotides
 9 obtained from human placenta, adult testes, fetal kidney, fetal
 10 brain, adult brain, adult brain and adult blood cDNA libraries.
 11 The polynucleotides and proteins are predicted to have biological
 12 activities which would make them suitable for treating, preventing or
 13 ameliorating medical conditions in humans and animals. Suggested
 14 activities include nutritional activity, cytokine and cell
 15 proliferation/differentiation activity, immune stimulation (e.g. as
 16 vaccines) or suppressing activity, haematopoiesis regulating
 17 activity, tissue growth activity, activin/inhibin activity,
 18 chemotactic/chemokinetic activity, haemostatic and thrombolytic
 19 activity, receptor/ligand activity, anti-inflammatory activity,
 20 cadherin/tumour invasion suppressor activity, and tumour inhibition
 21 activity. The polynucleotides are also stated to be useful for gene
 22 therapy. The sequences are identified by a secretory leader
 23 sequence motif in the polynucleotide and it is thought that the
 24 encoded proteins have biological activity by virtue of their secreted
 25 nature. This clone was designated AC222_1. A probe for this clone is
 26 described in X08698.
 27
 28 Sequence 1626 BP; 560 A; 327 C; 291 G; 443 T; 5 other;
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PS Disclosure; Page 18-19; 29pp; English.
 XX This sequence is the coding sequence for an example of the inhibitor of
 CC the invention (the encoded protein is not given in the specification).
 CC The inhibitor is a Kex2 proteinase family enzyme inhibitor with a
 CC molecular weight of 11,500. The inhibitor/protein (termed kexstatin) is
 CC expected to have pharmaceutical and pesticidal applications.
 XX
 SQ Sequence 2186 BP; 297 A; 827 C; 780 G; 282 T; 0 other;
 Query Match 0.6%; Score 18; DB 19; Length 2186;
 Best Local Similarity 100.0%; Pred. No. 55;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1652 ggtctctgggacccctggc 1669
 Db 1518 ggtctctgggacccctggc 1535
 RESULT 15
 X20537/c
 ID X20537 standard; DNA; 2450 BP.
 AC X20537;
 XX
 XX 05-MAY-1999 (first entry)
 XX
 DE Polynucleotide sequence from the genome of Treponema pallidum.
 XX
 KW Treponema pallidum infection; syphilis; Borrelia infection; animal;
 XX enzyme production; ds.
 OS Treponema pallidum.
 XX
 PN WO9859034-A2.
 PD 30-DEC-1998.
 XX
 PF 23-JUN-1998; 98WO-US13041.
 XX
 PR 24-JUN-1997; 97US-0050667.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Fraser CM;
 XX
 WPI; 1999-081273/07.
 XX
 PT New isolated Treponema pallidum nucleic acids - used to develop
 PT products for the detection, diagnosis, characterisation, prevention
 PT and therapy of T. pallidum infections, particularly syphilis
 XX
 PS Claim 1; Page 408-410; 1150pp; English.
 XX
 CC X20500-21243 represent polynucleotide sequences from the genome of
 CC Treponema pallidum. The sequences can be used for detection,
 CC diagnosis, characterisation, prevention and therapy for T. pallidum
 CC infections, particularly syphilis. They can also be used for detecting
 CC diseases related to Borrelia infections in animals, and for the
 CC production of biosynthetic products such as enzymes.
 XX
 SQ Sequence 2450 BP; 553 A; 643 C; 631 G; 617 T; 6 other;
 Query Match 0.6%; Score 18; DB 20; Length 2450;
 Best Local Similarity 100.0%; Pred. No. 55;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2305 tctgtcttggagacttcc 2322
 Db 941 TCTGCTTTGGAGACTTTC 924

(BIOM-) BIOMOLECULAR ENG RES INST.
 Kikuchi N, Oda K, Shibano Y;
 WPI; 1998-044337/05.
 Kex2 protease inhibitor protein - with potential pharmaceutical and
 pesticidal applications

```

RESULT 16
04699/c
V04699 standard; cDNA; 2605 BP.
V04699;
21-JUL-1998 (first entry)
Homo sapiens 20q13 amplicon cc43 transcript.
20q13 amplicon; chromosome 20; tumour; detection;
chromosomal abnormalities; probe; gene therapy; antisense inhibition;
treatment; age-related macular degeneration; retinitis pigmentation;
Leber's congenital amaurosis; ds.
Homo sapiens.
WO9802539-A1.
22-JAN-1998.
15-JUL-1997; 97WO-US12343.
17-JAN-1997; 97US-0785532.
15-JUL-1996; 96US-0680395.
16-OCT-1996; 96US-0731499.
(REGC ) UNIV CALIFORNIA.
Collins CC, Godfrey T, Gray JW, Hwang SI, Kowbel D;
Rommens J;
WPI; 1998-110587/10.
New sequences from the 20q13 amplicon - used for detecting
chromosomal abnormalities, particularly tumours, and for developing
products for treating diseases
Claim 1; Page 62; 91pp; English.
The sequence is that of a cDNA sequence cc43, which was isolated
from the 20q13 amplicon. It is expressed in normal tissues but not
been found in the breast cancer cell line. It can be used as a probe
for the detection of chromosomal abnormalities at 20q13. It and
other sequences isolated from the 20q13 amplicon are consistently
amplified in primary tumours. These sequences are useful as probes
or as probe targets for monitoring the relative copy number of
corresponding sequences from a biological sample such as tumour
cells. The sequences can also be used in therapeutic applications
for modulating the expression of the endogenous gene or the activity
of the gene product. Examples of therapeutic approaches include
antisense inhibition of gene expression, gene therapy, and monoclonal
antibodies that specifically bind the gene products. The products can
also be used in the treatment of other diseases, e.g. age-related
macular degeneration, Leber's congenital amaurosis and retinitis
pigmentation.
Sequence 2605 BP; 698 A; 625 C; 549 G; 732 T; 1 other;

Query Match 0.6%; Score 18; DB 19; Length 2605;
Best Local Similarity 100.0%; Pred. No. 55;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

448 ttccaaagtgtgtacttt 465
|||||
1421 TTCCAAAGTGTGACTTT 1404

RESULT 17
33895
233895 standard; cDNA; 2945 BP.

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XX AC Z33895;
XX DT 07-DEC-1999 (first entry)
XX DE Human PRO274 nucleotides sequence.
XX KW Human; PRO; EST; expressed sequence tag; PCR primer; hybridisation;
XX KW probe; blood coagulation disorder; cancer; cellular adhesion disorder;
XX KW secreted protein; transmembrane protein; ss.
XX OS Homo sapiens.
XX PN WO9946281-A2.
XX PD 16-SEP-1999.
XX PF 08-MAR-1999; 99WO-US05028.
XX PR 10-MAR-1998; 98US-0077450.
XX PR 11-MAR-1998; 98US-0077632.
XX PR 11-MAR-1998; 98US-0077641.
XX PR 11-MAR-1998; 98US-0077649.
XX PR 12-MAR-1998; 98US-0077791.
XX PR 13-MAR-1998; 98US-0078004.
XX PR 17-MAR-1998; 98US-0040220.
XX PR 20-MAR-1998; 98US-0078886.
XX PR 20-MAR-1998; 98US-0078910.
XX PR 20-MAR-1998; 98US-0078936.
XX PR 20-MAR-1998; 98US-0078939.
XX PR 25-MAR-1998; 98US-0079294.
XX PR 26-MAR-1998; 98US-0079656.
XX PR 27-MAR-1998; 98US-0079663.
XX PR 27-MAR-1998; 98US-0079664.
XX PR 27-MAR-1998; 98US-0079689.
XX PR 27-MAR-1998; 98US-0079728.
XX PR 27-MAR-1998; 98US-0079786.
XX PR 30-MAR-1998; 98US-0079920.
XX PR 30-MAR-1998; 98US-0079923.
XX PR 31-MAR-1998; 98US-0080105.
XX PR 31-MAR-1998; 98US-0080107.
XX PR 31-MAR-1998; 98US-0080165.
XX PR 01-APR-1998; 98US-0080194.
XX PR 01-APR-1998; 98US-0080327.
XX PR 01-APR-1998; 98US-0080328.
XX PR 01-APR-1998; 98US-0080333.
XX PR 01-APR-1998; 98US-0080334.
XX PR 08-APR-1998; 98US-0081049.
XX PR 08-APR-1998; 98US-0081070.
XX PR 08-APR-1998; 98US-0081071.
XX PR 09-APR-1998; 98US-0081195.
XX PR 09-APR-1998; 98US-0081203.
XX PR 09-APR-1998; 98US-0081229.
XX PR 15-APR-1998; 98US-0081817.
XX PR 15-APR-1998; 98US-0081838.
XX PR 15-APR-1998; 98US-0081952.
XX PR 15-APR-1998; 98US-0081955.
XX PR 21-APR-1998; 98US-0082568.
XX PR 21-APR-1998; 98US-0082569.
XX PR 22-APR-1998; 98US-0082700.
XX PR 22-APR-1998; 98US-0082704.
XX PR 22-APR-1998; 98US-0082804.
XX PR 23-APR-1998; 98US-0082767.
XX PR 23-APR-1998; 98US-0082796.
XX PR 27-APR-1998; 98US-0083336.
XX PR 28-APR-1998; 98US-0083322.
XX PR 29-APR-1998; 98US-0083392.
XX PR 29-APR-1998; 98US-0083495.
XX PR 29-APR-1998; 98US-0083496.
XX PR 29-APR-1998; 98US-0083499.
XX PR 29-APR-1998; 98US-0083500.
XX PR 29-APR-1998; 98US-0083545.
XX PR 29-APR-1998; 98US-0083554.

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CC Homo sapiens.
XX WO9846795-A1.
XX
XX 22-OCT-1998.
XX
XX 09-APR-1998; 98WO-US07027.
XX
XX 11-APR-1997; 97US-0843076.
XX
XX (BAYU ) BAYLOR COLLEGE MEDICINE.
XX (MAYO-) MAYO FOUNDATION.
XX
XX Slawin KM, Tindall DJ, Young CYF;
XX
XX WPI; 1998-594592/50.
XX
XX Detection of human kallikrein 2 RNA - by reverse transcription and
XX amplification by PCR, for detecting, monitoring and staging of
XX prostate cancer
XX
XX Disclosure: Page 78-80; 90pp; English.
XX
XX The present invention describes a diagnostic method for detecting human
XX kallikrein 2 (hk2) DNA. The method comprises: (a) contacting DNA obtained
XX by reverse transcription (RT) of RNA from a human physiological sample
XX which comprises cells suspected of containing hk2 RNA with at least 2
XX oligonucleotides to amplify the DNA by PCR to yield amplified hk2 DNA,
XX where the conditions amplify the DNA obtained by RT of RNA from at least
XX one cell containing hk2 in a sample which comprises at least 107 to 109
XX cells; and (b) detecting the presence of the amplified hk2 DNA. The
XX method can be used for detecting, monitoring the progression of and
XX pathologically staging prostate cancer. The present sequence represents
XX the coding strand of native genomic hk2.
XX
XX Sequence 6139 BP; 1233 A; 1875 C; 1595 G; 1436 T; 0 other;

Query Match 0.6%; Score 18; DB 19; Length 6139;
Best Local Similarity 100.0%; Pred. No. 54;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1048 cagatgccccctggcct 1065
D 3041 cagatgccccctggcct 3058
|||||
RESULT 20
123903/C
D Z23903 standard; DNA; 49999 BP.
XX
XX Z23903;
XX
XX 25-JAN-2000 (first entry)
XX
XX Human LOBO homologue genomic DNA fragment 5.
XX
XX LOBO; long bones; bone development; bone extension; skull; osteopathic;
XX diagnostic; pharmaceutical; gene therapy; transgenic animal; disease;
XX spondyloepiphyseal dysplasia; achondroplasia; human; ds.
XX
XX Homo sapiens.
XX
XX WO9950284-A2.
XX
XX 07-OCT-1999.
XX
XX 26-MAR-1999; 99WO-EP02055.
XX
XX 27-MAR-1998; 98DE-1013799.
XX
XX (ROSE/) ROSENTHAL A.

PI Rosenthal A, Rump A, Hess J, Aigner T, Wirth T;
XX
XX WPI; 1999-601320/51.
XX
XX Nucleic acids encoding proteins which influence bone development,
XX useful for treating and studying bone disorders -
XX
XX Example 3; Page 328-356; 391pp; German.
XX
XX This invention describes novel nucleic acids (I; designated LOBO (long
XX bones)) encoding proteins influencing bone development in mammals. The
XX proteins of the invention reduce and/or inactivate bone extension (i.e.
XX development), with exception of the skull and have osteopathic activity.
XX The nucleic acid molecules, proteins and antibodies can be used in
XX diagnostic or pharmaceutical compounds e.g. for gene therapy. The methods
XX and nucleic acid molecules, etc. are useful for production of transgenic
XX animals, especially a transgenic mouse for the study of diseases
XX associated with bone development, e.g. spondyloepiphyseal dysplasia and
XX achondroplasia. This sequence encodes a human LOBO protein described
XX in the method of the invention.
XX
XX Sequence 49999 BP; 10983 A; 13723 C; 13439 G; 11854 T; 0 other;

Query Match 0.6%; Score 18; DB 20; Length 49999;
Best Local Similarity 100.0%; Pred. No. 52;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2740 gggccagagagctggcct 2757
D 14820 GGGCCAGGAGGCTGCCAC 14803
|||||
RESULT 21
Q54684
ID Q54684 standard; DNA; 21 BP.
XX
XX Q54684;
XX
XX 05-JUL-1994 (first entry)
XX
XX C kappa exon primer.
XX
XX Germ line; stem cell; blastocyst; implantation; embryonic;
XX electroporation; kappa; constant; exon; ss.
XX
XX Synthetic.
XX
XX DE4228162-C.
XX
XX 13-JAN-1994.
XX
XX 25-AUG-1992; 92DE-4228162.
XX
XX 25-AUG-1992; 92DE-4228162.
XX
XX (RAJE/) RAJEWSKY K.
XX (KOEL-) KOELNER VER FORDERUNG IMMUNOLOGIE.
XX
XX Rajewsky K, Zou Y;
XX
XX WPI; 1994-008862/02.
XX
XX Homologous replacement of gene in mammalian germ line - by
XX transfecting embryonic stem cell with labelled recombination
XX vehicle, selection and incorporating into blastocyst(s) for
XX implantation, esp. for prodn. of humanised antibodies in mice
XX
XX Example; Column 4; 7pp; German.
XX
XX DSM 7211 is made by (1) inserting a fragment of pc-2 (contg.
XX the C kappa exon) into pTZ-19(R); (2) inserting a 1.2 kb
XX fragment of pBHC kappa (contg. the intron enhancer element);

```


Human 5' EST isolated from a cDNA library SEQ ID NO:402.

Human; 5' EST; expressed sequence tag; secreted protein; diagnosis; gene therapy; chromosome mapping; upstream regulatory sequence; forensic; location; development; protein synthesis; stability; regulation; identification; ss.

Homo sapiens.

WO9953051-A2.

21-OCT-1999.

09-APR-1999; 99WO-IB00712.

09-APR-1998; 98US-0057719.

28-APR-1998; 98US-0069047.

(GEST) GENSET.

Dumas Milne Edwards J, Duclert A, Giordano J;

WPI; 2000-038446/03.

P-PSDB; Y65029.

Novel secreted protein 5' expressed sequence tag sequences used in diagnostic, forensic, gene therapy, and chromosome mapping procedures

Claim 1; Page 382; 837pp; English.

Z42265 to Z43075 represent novel 5' expressed sequence tag (EST) sequences, corresponding to human secreted proteins. Y64551 to Y65438 represent the EST-related proteins corresponding to Z42265 to Z43052. The 5' ESTs can be used for producing secreted human gene products. They can be used to identify and isolate 5' untranslated regions (UTRs) and upstream regulatory regions which control the location, development stage, rate, and quantity of protein synthesis, as well as stability of mRNA. The ESTs are also useful as probes for chromosome mapping, and to obtain full length cDNA clones. The ESTs can also be used in forensic procedures to identify individuals, or in diagnostic procedures to identify individuals having genetic diseases resulting from abnormal gene expression. The products may also be used in gene therapy protocols. The nucleic acids encoding signal peptides can be used for directing extracellular secretion of a polypeptide or the insertion of a polypeptide into a membrane, or importing a polypeptide into a cell. The proteins encoded by the EST sequences may be useful in treating a variety of human conditions. Secreted proteins have therapeutic value, and the identification of new secreted proteins is valuable. Z42249 to Z42264 and Y64644 to Y64650 represent sequences used in the exemplification of the present invention.

Sequence 330 BP; 76 A; 102 C; 86 G; 65 T; 1 other;

Query Match 0.6%; Score 17; DB 21; Length 330;

Best Local Similarity 100.0%; Pred. No. 1.8e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1219 acctcatccaccggac 1235

|||||

48 ACCTCATCCACCGGAC 32;

RESULT 27

X24993/c

X24993 standard; cDNA; 332 BP.

X24993;

05-JUL-1999 (first entry)

Murine Bcl-2 interacting mediator of cell death Bim-S cDNA.

Bim-S; Bcl-2 interacting mediator of cell death; apoptosis; cell cycle; mouse; cancer; autoimmune disease; splice variant; degenerative disease; therapy; contraceptive; isoform; ss.

Mus musculus.

WO9914321-A1.

25-MAR-1999.

17-SEP-1998; 98WO-AU00772.

24-SEP-1997; 97AU-0009373.

17-SEP-1997; 97AU-0009263.

(HALL-) HALL INST MEDICAL RES WALTER & ELIZA.

Adams J, Cory S, Huang DCS, O'Connor L, O'Reilly L;

Puthalakath H, Strasser A;

WPI; 1999-244030/20.

P-PSDB; W98154.

New isolated member of the Bcl-2 family, Bim used in, e.g. cancer treatment

Claim 3; Page 92; 145pp; English.

The present sequence encodes the short form (S) of murine Bim, or Bcl-2 interacting mediator of cell death (see W98154), a novel member of the Bcl-2 family that is capable of inducing cell death (apoptosis) and which acts as a death-ligand for certain members of the pro-survival Bcl-2 family. Bim is a BH3-only protein, as the only Bcl-2 homology region which it encompasses is BH3. It is the only BH3-only protein for which splice variants exist. These result in the expression of a variety of isoforms, i.e. Bim-S, Bim-L and Bim-EL (see W98154-56). cDNAs encoding these murine Bim isoforms were obtained from a T lymphoma cDNA library using human recombinant Bcl-2 protein. The murine Bim gene has been mapped to chromosome 2 at bands F3-G. Human Bim-L and Bim-EL isoforms have also been identified (see W98157-58). Binding the dynein light chain was shown to regulate the pro-apoptotic activity of Bim. Bim-S, the splice variant which does not bind to dynein light chain, is a much more potent killer than either Bim-L or Bim-EL. The invention provides variants (see W98159-68) of murine and human Bim-L or Bim-EL that cannot bind, couple or otherwise associate with a dynein light chain. The identification of Bim permits the identification and rational design of a range of products for use in therapy, diagnosis, antibody generation and involving modulation of physiological cell death. These therapeutic molecules may act as either antagonists or agonists of Bim's function and will be useful in cancer, autoimmune or degenerative disease therapy. Increased Bim expression or Bim activity is useful, e.g. for treatment or prophylaxis in conditions such as cancer and deletion of autoreactive lymphocytes in autoimmune disease. Decreased Bim expression of Bim activity is useful in regulating inhibition or prevention of cell death or degeneration such as under cytotoxic conditions during e.g. gamma-irradiation and chemotherapy or during HIV/AIDS or other viral infections, ischemia, myocardial infarction, hypoxia, degenerative diseases or for prolonging the survival of cells being transplanted for treatment of disease. Since Bim is expressed in germ cells, modulating Bim expression or Bim activity is useful, e.g. as a contraceptive or method of sterilization by preventing generation of fertile sperm.

Sequence 332 BP; 87 A; 85 C; 91 G; 69 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 332;

Best Local Similarity 100.0%; Pred. No. 1.8e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

654 aggcctctcagcaggct 670
|||||
75 AGGCTCTCAGCAGGCT 59

RESULT 28

T24629 standard; cDNA to mRNA; 347 BP.

T24629;

07-OCT-1996 (first entry)

Human gene signature HUMGS06689.

Gene signature: messenger RNA; mRNA; relative abundance; frequency;
human; cloning; mapping; non-biased library; diagnosis; detection;
cell typing; abnormal cell function; ss.

Homo sapiens.

W09514772-A1.

01-JUN-1995.

11-NOV-1994; 94WO-JP01916.

12-NOV-1993; 93JP-0355504.

(MATS/) MATSUBARA K.
(OKUB/) OKUBO K.

Matsubara K, Okubo K;

WPI: 1995-206931/27.

Identifying gene signatures in 3'-directed human cDNA library - e.g.
for diagnosis of abnormal cell function, by preparing cDNA that
reflects relative abundance of corresp. mRNA in specific human
tissues

Claim 1; Page 1655; 2245pp; Japanese.

A single-stranded DNA (or its complementary strand or the corresp.
double-stranded DNA) which comprises one of the 7837 "GS" sequences
given in T19001-T26837 and which is able to hybridise to part of
human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
sequences were obtained from 3'-directed cDNA libraries prepared
from various human tissues; synthesis of cDNA was initiated from the
3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
untranslated sequence is unique to a particular mRNA species, almost
all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
is constructed so as to reflect accurately the relative abundance of
different mRNAs in the particular tissue from which it was derived.
The appearance frequency of a given GS in a cDNA library can be
determined (esp. using primers and probes derived from the GS
sequences) as a means of diagnosing abnormal cell function or for
recognising different cell types.

Sequence 347 BP; 61 A; 102 C; 108 G; 75 T; 1 other;

Query Match 0.6%; Score 17; DB 16; Length 347;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 1336 ccaggaggagtgagtcag 1352

|||||

15 ccaggaggaggagtgagtcag 31

RESULT 29

X24994/c

X41005 standard; cDNA; 369 BP.

X41005;

18-JUN-1999 (first entry)

Human secreted protein 5' EST SEQ ID NO: 217.

Human; secreted protein; EST; expressed sequence tag; diagnosis;

forensic; gene therapy; chromosome mapping; signal peptide;

upstream regulatory sequence; cytokine activity; cell proliferation;

differentiation; haematopoiesis regulation; tissue growth regulation;

reproductive hormone regulation; chemotactic; chemokinetic; haemostatic;

thrombolytic; anti-inflammatory; tumour inhibition; ds.

Homo sapiens.

W09906554-A2.

11-FEB-1999.

31-JUL-1998; 98WO-IB01238.

01-AUG-1997; 97US-0905134.

(GEST) GENSET.

Duciert A, Dumas Milne Edwards J, Lacroix B;

WPI: 1999-153784/13.

P-PSDB; Y12172.

New nucleic acids encoding human secreted proteins - obtained from
cDNA libraries prepared from kidney, fetal kidney, dystrophic
muscle, muscle and heart tissue

Claim 1; Page 314-315; 622pp; English.

X40826 to X41093 represent 5' expressed sequence tags (ESTs) for human
secreted proteins, and encode the proteins given in Y01602 and
Y11994 to Y12260, respectively. The proteins given represent the signal
peptide and an N-terminal fragment of a secreted protein. The nucleic
acid sequences can be used for producing secreted human gene products.
They can also be used to develop products for diagnosis and therapy.

The proteins obtained may have cytokine activity, cell

proliferation/differentiation activity, haematopoiesis regulating

activity, tissue growth regulating activity, reproductive hormone

regulating activity, chemotactic/chemokinetic activity, haemostatic and

thrombolytic activity, receptor/ligand activity, anti-inflammatory

activity, tumour inhibition activity or other activities. The products

can be used in forensic, gene therapy and chromosome mapping procedures.

The sequences can also be used for obtaining corresponding promoter

sequences. The nucleic acids encoding the signal peptide can be used

for directing extracellular secretion of a polypeptide or the insertion

of a polypeptide into a membrane, or importing a polypeptide into

a cell.

Sequence 369 BP; 57 A; 128 C; 98 G; 84 T; 2 other;

Query Match 0.6%; Score 17; DB 20; Length 369;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2501 gccacagcccaagaaggg 2517

|||||

Db 320 GCACAGCCCAAGAAGG 304

RESULT 30

X24994/c

ID X24994 standard; cDNA; 422 BP.

XX

CC XZ4994;

XX 05-JUL-1999 (first entry)

XX Murine Bcl-2 interacting mediator of cell death Bim-L cDNA.

XX Bim-L; Bcl-2 interacting mediator of cell death; apoptosis;

XX cell cycle; mouse; cancer; autoimmune disease;

XX degenerative disease; therapy; contraceptive; splice variant;

XX isoform; ss.

XX Mus musculus.

XX WO9914321-A1.

XX 25-MAR-1999.

XX 17-SEP-1998; 98WO-AU000772.

XX 24-SEP-1997; 97AU-0009373.

XX 17-SEP-1997; 97AU-0009263.

XX (HALL-) HALL INST MEDICAL RES WALTER & ELIZA.

XX Adams J, Cory S, Huang DCS, O'Connor L, O'Reilly L;

XX Puthalakath H, Strasser A;

XX WPI; 1999-244030/20.

XX P-ESDB; W98155.

XX New isolated member of the Bcl-2 family, Bim used in, e.g. cancer

XX treatment

XX Claim 3; Page 94-95; 145pp; English.

XX The present sequence encodes the long form (L) of murine Bim, or

XX Bcl-2 interacting mediator of cell death (see W98155), a novel

XX member of the Bcl-2 family that is capable of inducing cell death

XX (apoptosis) and which acts as a 'death-ligand' for certain members

XX of the pro-survival Bcl-2 family. Bim is a BH3-only protein, as the

XX only BH3-only protein region which it encompasses is BH3. It is the

XX result in the expression of a variety of isoforms, i.e. Bim-S,

XX Bim-L and Bim-EL (see W98154-56). cDNAs encoding these murine Bim

XX isoforms were obtained from a T lymphoma cDNA library using human

XX recombinant Bcl-2 protein. The murine Bim gene has been mapped to

XX chromosome 2 at bands F3-G. Human Bim-L and Bim-EL isoforms have

XX also been identified (see W98157-58). Binding the dynein light

XX chain was shown to regulate the pro-apoptotic activity of Bim.

XX Bim-S, the splice variant which does not bind to dynein light

XX chain, is a much more potent killer than either Bim-L or Bim-EL.

XX The invention provides variants (see W98159-68) of murine and human

XX Bim-L or Bim-EL that cannot bind, couple or otherwise associate

XX with a dynein light chain. The identification of Bim permits the

XX identification and rational design of a range of products for use

XX in therapy, diagnosis, antibody generation and involving modulation

XX of physiological cell death. These therapeutic molecules may act

XX as either antagonists or agonists of Bim's function and will be

XX useful in cancer, autoimmune or degenerative disease therapy.

XX Increased Bim expression or Bim activity is useful, e.g. for

XX treatment or prophylaxis in conditions such as cancer and deletion

XX of autoreactive lymphocytes in autoimmune disease. Decreased Bim

XX expression of Bim activity is useful in regulating inhibition or

XX prevention of cell death or degeneration such as under cytotoxic

XX conditions during e.g. gamma-irradiation and chemotherapy or during

XX HIV/AIDS or other viral infections, ischemia, myocardial infarction,

XX hypoxia, degenerative diseases or for prolonging the survival of

XX cells being transplanted for treatment of disease. Since Bim is

XX expressed in germ cells, modulating Bim expression or Bim activity

XX is useful, e.g. as a contraceptive or method of sterilization by

XX preventing generation of fertile sperm.

XX Sequence 422 BP; 112A; 116 C; 109 G; 85 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 422;

Best Local Similarity 100.0%; Pred. No. 1.8e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 654 aggcctctcagcagct 670

|||||

DB 75 AGGCTCTCAGCAGCT 59

RESULT 31

A32069/C

ID A32069 standard; DNA; 426 BP.

XX AC A32069;

DT 05-JUL-2000 (first entry)

DE Plant microsatellite marker #1030.

XX Plant microsatellite sequence; core repeat sequence; detection; probe;

KW DNA polymorphism; genome mapping; physical mapping; fingerprinting;

XX variety identification; genetic variability evaluation; primer; ss.

OS Pinus radiata.

XX WO9967421-A1.

XX 29-DEC-1999.

XX 25-JUN-1999; 99WO-NZ00092.

XX 25-JUN-1998; 98US-0105307.

XX (GENE-) GENESIS RES & DEV CORP LTD & FLETCHER.

XX (FLET-) FLETCHER CHALLENGE FORESTS LTD.

XX Havukkala IJ, Bloksberg LN, Glenn M;

XX WPI; 2000-116958/10.

XX New plant microsatellite markers and associated flanking species for

XX the detection of polymorphic genetic markers -

XX Claim 1; Page 379; 392pp; English.

XX Sequences A31040-A32093 represent novel plant microsatellite sequences

XX and associated flanking species. The sequences comprise a central core

XX repeat sequence, especially selected from the sequences A32094-A32095

XX with left and right flanking sequences. The polynucleotide sequences

XX can be used in the detection of DNA polymorphisms, in genome mapping,

XX in physical mapping, in positional cloning of genes, in variety

XX identification and in evaluation of genetic variability within and

XX between plant tissues, populations, cultivars, species and species

XX groups. They may also be used to design hybridization probes for

XX oligonucleotide fingerprinting and library screening and to design

XX primers for microsatellite-primed PCR. Microsatellite markers are

XX useful to locate specific economically useful genes in plant genomes.

XX Sequence 426 BP; 124 A; 81 C; 71 G; 150 T; 0 other;

Query Match 0.6%; Score 17; DB 21; Length 426;

Best Local Similarity 100.0%; Pred. No. 1.8e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2021 tggctggaaagtgtct 2037

|||||

DB 257 TGGCTGGAAGTGTCT 241

RESULT 32

32052/C
 C A32052 standard; DNA; 431 BP.
 XX
 A32052;
 XX
 05-JUL-2000 (first entry)
 XX
 Plant microsatellite marker #1013.
 XX
 Plant microsatellite sequence; core repeat sequence; detection; probe;
 XX DNA polymorphism; genome mapping; physical mapping; fingerprinting;
 XX variety identification; genetic variability evaluation; primer; ss.
 XX
 Pinus radiata.
 XX
 WO9967421-A1.
 XX
 29-DEC-1999.
 XX
 25-JUN-1999; 99WO-NZ00092.
 XX
 25-JUN-1998; 98US-0105307.
 XX
 (GENE-) GENESIS RES & DEV CORP LTD & FLETCHER.
 XX (FLET-) FLETCHER CHALLENGE FORESTS LTD.
 XX
 Havukkala IJ, Bloksberg LN, Glenn M;
 XX
 WPI; 2000-116958/10.
 XX
 New plant microsatellite markers and associated flanking species for
 XX the detection of polymorphic genetic markers -
 XX
 Claim 1; Page 374; 392pp; English.
 XX
 Sequences A31040-A32093 represent novel plant microsatellite sequences
 XX and associated flanking species. The sequences comprise a central core
 XX repeat sequence, especially selected from the sequences A32094-A32096
 XX with left and right flanking sequences. The polynucleotide sequences
 XX can be used in the detection of DNA polymorphisms, in genome mapping,
 XX in physical mapping, in positional cloning of genes, in variety
 XX identification and in evaluation of genetic variability within and
 XX between plant tissues, populations, cultivars, species and species
 XX groups. They may also be used to design hybridization probes for
 XX oligonucleotide fingerprinting and library screening and to design
 XX primers for microsatellite-primed PCR. Microsatellite markers are
 XX useful to locate specific economically useful genes in plant genomes.
 XX
 Sequence 431 BP; 129 A; 78 C; 71 G; 152 T; 1 other;
 XX
 Query Match 0.6%; Score 17; DB 21; Length 431;
 Best Local Similarity 100.0%; Pred. No. 1.8e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 2021 tggctggaagtgtct 2037
 ||||||||||||||||
 262 TGCTGGAAAGTGTCT 246
 XX
 RESULT 33
 V29360/C
 V29360 standard; DNA; 559 BP.
 XX
 V29360;
 XX
 31-JUL-1998 (first entry)
 XX
 Calcium ion channel alpha1 subunit exons 33, 34/intron; partial sequence.
 XX
 Calcium ion channel alpha1 subunit; human; episodic ataxia type 2;
 XX familial hemiplegic migraine; FHM; EA-2; treatment; diagnosis;
 XX exon; intron; ss.

XX OS Homo sapiens.
 XX
 XX FH Key
 FT intron Location/Qualifiers
 FT 1..156
 FT /tag= a
 FT /number= 32
 FT /note= "partial sequence"
 FT 157..222
 FT /tag= b
 FT /number= 33
 FT 223..394
 FT /tag= c
 FT /number= 33
 FT 395..509
 FT /tag= d
 FT /number= 34
 FT 510..559
 FT /tag= e
 FT /number= 34
 FT /note= "partial sequence"
 FT
 XX EP834561-A1.
 PN
 08-APR-1998.
 PD
 XX
 XX 27-SEP-1996; 96EP-0202707.
 PF
 XX 27-SEP-1996; 96EP-0202707.
 PR
 XX (UYLE-) RIJKSUNIV LEIDEN.
 PA
 XX Ferrari MD, Frants RRIE, Ophoff RA, Terwindt GM;
 PI
 XX WPI; 1998-195461/18.
 DR
 XX
 XX New human nucleic acid associated with migraine and episodic ataxia
 PT type 2 - useful for diagnosis and development of specific treatments
 PT
 XX Disclosure; Fig 1; 157pp; English.
 PS
 XX Sequences shown in V29330 to V29371 represent the 47 exons and flanking
 CC intronic sequences containing the complete coding region of the human
 CC calcium ion channel alpha 1 subunit gene and part of untranslated
 CC sequences. The channel is related to familial hemiplegic migraine (FHM)
 CC and/or episodic ataxia type 2 (EA-2) and is derived from, related to or
 CC associated with a gene present in humans on chromosome 19p13.1-13.2. The
 CC encoding gene can be used to localise or identify genes related to
 CC episodic neurological disorders, specifically migraine, FHM or EA-2, but
 CC also epilepsy. The isolated or a recombinant nucleic acid can also be
 CC used to distinguish between alleles of the corresponding gene. Cells and
 CC animals containing recombinant expression vectors comprising the nucleic
 CC acid can be useful in study, development and treatment of migraine, FHM,
 CC EA-2 and epilepsy. Proteins or peptides encoded by the nucleic acid and
 CC natural or synthetic antibodies against the proteins can be used to
 CC diagnose FHM, EA-2, migraine and other neurological conditions associated
 CC with cation channel dysfunction.
 CC
 XX Sequence 559 BP; 119 A; 157 C; 160 G; 120 T; 3 other;
 XX
 Query Match 0.6%; Score 17; DB 19; Length 559;
 Best Local Similarity 100.0%; Pred. No. 1.8e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 QY 1207 aaaccagctcaacctc 1223
 ||||||||||||||||
 DB 339 AAACCCAGCTCAACCTC 323
 XX
 RESULT 34
 ID X24995/c
 X24995 standard; cDNA; 590 BP.

XX X24995;
XX 05-JUL-1999 (first entry)
XX Murine Bcl-2 interacting mediator of cell death Bim-EL CDNA.
XX Bim-EL; Bcl-2 interacting mediator of cell death; apoptosis;
XX cell cycle; mouse; cancer; autoimmune disease;
XX degenerative disease; therapy; contraceptive; splice variant;
XX isoform; ss.
XX
XX Mus musculus.
XX WO9914321-A1.
XX
XX 25-MAR-1999.
XX
XX 17-SEP-1998; 98WO-AU00772.
XX
XX 24-SEP-1997; 97AU-0009373.
XX 17-SEP-1997; 97AU-0009263.
XX (HALL-) HALL INST MEDICAL RES WALTER & ELIZA.
XX Adams J, Cory S, Huang DCS, O'Connor L, O'Reilly L;
XX Puthalakath H, Strasser A;
XX WPI; 1999-244030/20.
XX P-PSDB; W98156.
XX New isolated member of the Bcl-2 family, Bim used in, e.g. cancer
XX treatment
XX
XX Claim 3; Page 96-97; 145pp; English.
XX
XX The present sequence encodes the extra long form (EL) of murine Bim,
XX or Bcl-2 interacting mediator of cell death (see W98156), a novel
XX member of the Bcl-2 family that is capable of inducing cell death
XX (apoptosis) and which acts as a 'death-ligand' for certain members
XX of the pro-survival Bcl-2 family. Bim is a BH3-only protein, as the
XX only Bcl-2 homology region which it encompasses is BH3. It is the
XX only BH3-only protein for which splice variants exist. These
XX result in the expression of a variety of isoforms, i.e. Bim-S,
XX Bim-L and Bim-EL (see W98154-56). cDNAs encoding these murine Bim
XX isoforms were obtained from a T lymphoma cDNA library using human
XX recombinant Bcl-2 protein. The murine Bim gene has been mapped to
XX chromosome 2 at bands F3-G. Human Bim-L and Bim-EL isoforms have
XX also been identified (see W98157-58). Binding the dynein light
XX chain was shown to regulate the pro-apoptotic activity of Bim.
XX Bim-S, the splice variant which does not bind to dynein light
XX chain, is a much more potent killer than either Bim-L or Bim-EL.
XX The invention provides variants (see W98159-68) of murine and human
XX Bim-L or Bim-EL that cannot bind, couple or otherwise associate
XX with a dynein light chain. The identification of Bim permits the
XX identification and rational design of a range of products for use
XX in therapy, diagnosis, antibody generation and involving modulation
XX of physiological cell death. These therapeutic molecules may act
XX as either antagonists or agonists of Bim's function and will be
XX useful in cancer, autoimmune or degenerative disease therapy.
XX Increased Bim expression or Bim activity is useful, e.g. for
XX treatment or prophylaxis in conditions such as cancer and deletion
XX of autoreactive lymphocytes in autoimmune disease. Decreased Bim
XX expression of Bim activity is useful in regulating inhibition or
XX prevention of cell death or degeneration such as under cytotoxic
XX conditions during e.g. gamma-irradiation and chemotherapy or during
XX HIV/AIDS or other viral infections, ischemia, myocardial infarction,
XX hypoxia, degenerative diseases or for prolonging the survival of
XX cells being transplanted for treatment of disease. Since Bim is
XX expressed in germ cells, modulating Bim expression or Bim activity
XX is useful, e.g. as a contraceptive or method of sterilization by
XX preventing generation of fertile sperm.

SQ Sequence 590 BP; 137 A; 178 C; 150 G; 125 T; 0 other;
Query Match 0.6%; Score 17; DB 20; Length 590;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 654 aggcctctcagcaggct 670
Db ||||||||||||||||
75 AGGCCTCTCAGCAGGCT 59
RESULT 35
V87198/C
ID V87198 standard; CDNA; 605 BP.
XX
XX AC V87198;
XX
XX DT 27-APR-1999 (first entry)
XX
XX DE EST clone BN180.
XX
XX KW Expressed sequence tag; secreted protein; haematopoiesis regulator;
XX tissue growth; activin; inhibin; tumour invasion suppressor; EST; human;
XX chemotaxis; chemokinesis; haemostasis; gene therapy; thrombolysis;
XX receptor; ligand; anti-inflammatory; tumour inhibitor; ds.
XX
XX OS Homo sapiens.
XX
XX PN WO9845435-A2.
XX
XX PD 15-OCT-1998.
XX
XX PF 10-APR-1998; 98WO-US06954.
XX
XX PR 10-APR-1997; 97US-0835913.
XX
XX PA (GENY) GENETICS INST INC.
XX
XX PI Agostino MJ, Jacobs K, Lavallie ER, McCoy JM, Merberg D;
XX PI Racie LA, Spaulding V, Treacy M;
XX WPI; 1999-070076/06.
XX
XX PT New polynucleotides encoding human secreted proteins - derived from
XX e.g. human blood, kidney, foetal lung, placenta, testes, brain,
XX ovary, pituitary, retina and colon cDNA libraries
XX
XX PS Claim 1; Page 486; 633pp; English.
XX
XX CC This sequence represents an expressed sequence tag (EST), and is a
XX polynucleotide of the invention. The polynucleotides of the invention are
XX all secreted EST sequences isolated from a variety of human tissue
XX sources. The EST sequences and proteins encoded by them are predicted to
XX have useful biological activities which would make them suitable for
XX treating, preventing or ameliorating medical conditions in humans and
XX animals, although no supporting data is given. Suggested activities
XX include nutritional activity, immune stimulating or suppressing activity,
XX haematopoiesis regulating activity, tissue growth activity,
XX activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
XX and thrombolytic activity, receptor/ligand activity, anti-inflammatory
XX activity, cadherin/tumour invasion suppressor activity, tumour inhibition
XX activity. The EST sequences are also stated to be useful for gene
XX therapy.
XX
XX SQ Sequence 605 BP; 149 A; 126 C; 194 G; 136 T; 0 other;
Query Match 0.6%; Score 17; DB 20; Length 605;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 31 gttttggggagacggg 47

212 GTTTGCTGGAGACGGG 196

RESULT 36
24979/C
X24979 standard; cDNA; 791 BP.

X24979;

05-JUL-1999 (first entry)

Clone GJ156 encoding TRAIN-R secreted form C-terminus.

TRAIN-R; receptor; human; tumour necrosis factor receptor;
agonist; antagonist; cancer; immunological disease; therapy;
cytostatic; ss.

Homo sapiens.

Key Location/Qualifiers

Intron 1..350

/*tag= a

351..790

/*tag= b

352..444

/*tag= c

/partial

/product= "TRAIN-R secreted form C-terminus"

45..790

/*tag= d

3'UTR

WO9913078-A1.

18-MAR-1999.

11-SEP-1998; 98WO-0519030.

06-MAY-1998; 98US-0084422.

12-SEP-1997; 97US-0058631.

(BIOJ) BIOGEN INC.

Hession C, Tschopp J;

WPI; 1999-229238/19.

P-PSDB; W98147.

New cysteine-rich tumor necrosis factor receptor

Claim 1; Page 28; 30pp; English.

The present sequence includes an exon encoding the C-terminus (see W98147) of a soluble form of a novel human cysteine-rich tumour necrosis factor receptor family member termed TRAIN-R. It comprises clone GJ156, obtained from a Clontech human adult lung cDNA library. The encoded 30-amino acid C-terminal peptide is identical to amino acids 121-149 of the composite TRAIN-R protein given in W98146 and to amino acids 121-150 of the C-terminus of murine TRAIN-R short form (secreted protein, see W98144). The soluble protein is expected to inhibit signalling by the full-length TRAIN-R. Human TRAIN-R is expressed at low levels in every tissue and cell line tested thus far, with higher expression detected in heart, prostate, ovary, testis, peripheral blood lymphocytes, thyroid and adrenal gland. Cell death can be induced by administering an agent capable of inhibiting the binding of TRAIN-R to its ligand. A claimed method of treating, or reducing, the advancement, severity or effects of an immunological disease in a mammal comprises administering a pharmaceutical composition which comprises a TRAIN-R blocking agent, e.g. soluble TRAIN-R. TRAIN-R can be fused to an immunoglobulin to produce a fusion protein which may be targeted to various sites. It can be used in binding assays, and to identify antagonists and agonists. Anti-TRAIN-R antibodies can be used to reduce the

CC severity of an immune response or to treat cancer. TRAIN-R
CC blocking agents can also be used to reduce the severity or effects
CC of an immunological disease (all claimed).

SQ Sequence 791 BP; 202 A; 189 C; 165 G; 235 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 791;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2130 gaagaggaagcagtgga 2146

Db 203 GAAGAGGAGCAGTGGA 187

RESULT 37

Z91823

ID Z91823 standard; DNA; 855 BP.

XX AC Z91823;

XX DT 02-JUN-2000 (first entry)

XX DE Streptococcus pneumoniae DNA sequence ID33.

XX KW Streptococcus pneumoniae infection; immunogen; antigen; diagnosis; AIDS;
KW bacterial pneumonia; asplenia; heart disease; lung disease; alcoholism;
KW kidney disease; diabetes; immunosuppressive disorder; otitis media;
KW pneumococcal septicaemia; sinusitis; meningitis; therapy; ss.

XX OS Streptococcus pneumoniae.

XX PN WO200006738-A2.

XX PD 10-FEB-2000.

XX PF 27-JUL-1999; 99WO-GB02452.

XX PR 27-JUL-1998; 98GB-0016336.

XX PR 19-MAR-1999; 99US-0125329.

XX PA (MICK-) MICROBIAL TECHNIKS LTD.

XX PI Le Page RWF, Wells JM, Hanniffy SB, Hansbro PM;

XX DR WPI; 2000-195301/17.

XX DR P-PSDB; Y81727.

XX PT Streptococcal proteins and polynucleotides useful for diagnosis,
XX treatment and prophylaxis of bacterial infections

PS Claim 2; Page 47-48; 76pp; English.

XX This sequence encodes a Streptococcus pneumoniae protein of the
XX invention. The proteins (or their homologues, derivatives and/or
XX fragments) are useful as immunogens or antigens. Immunogenic or antigenic
XX compositions comprising the proteins are useful as vaccines and also in
XX diagnostic assays. The sequences are useful for the detection or
XX diagnosis of S. pneumoniae infection, by contacting a sample to be tested
XX with them. Agents capable of antagonising, inhibiting or interfering with
XX the function of expression of the protein or polypeptide are useful in
XX medical compositions in the treatment or prophylaxis of S. pneumoniae
XX infection. As the sequences can be used to treat S. pneumoniae infection,
XX they can be used to treat bacterial pneumonia, which has high rates in
XX young children, the elderly, and in patients with predisposing conditions
XX such as asplenia, heart, lung and kidney disease, diabetes, alcoholism,
XX or with immunosuppressive disorders, especially AIDS. They can also be
XX used to treat pneumococcal septicaemia, otitis media, sinusitis, and
XX meningitis.

XX SQ Sequence 855 BP; 235 A; 173 C; 202 G; 245 T; 0 other;


```

Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggcaccctgg 1668
   |||||
DB 203 ggtcctgggcaccctgg 219

RESULT 39
Q67866
ID Q67866 standard; DNA; 1084 BP.
XX Q67866;
XX
XX 22-MAR-1995 (first entry)
XX
XX H6/MAGE-1 expression cassette from pMAW037.
XX
XX Polymerase chain reaction; primer; amplify; NTVAC; ALVAC; recombinant;
XX human; MAGE-1; melanoma-associated antigen; MZ2-E; testis; PTZ18RWAGE1;
XX primary melanoma tumour cell; melanoma-derived cell line; tumour;
XX poxvirus; antigenic response; immunological response; pathogen; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX FH misc_feature 1..51
XX FT /*tag= a
XX FT /note= "Flanking sequence"
XX FT promoter 52..178
XX FT /*tag= b
XX FT /note= "Vaccinia H6 promoter"
XX FT CDS 179..1009
XX FT /*tag= c
XX FT /product= MAGE-1
XX FT misc_feature 1010..1084
XX FT /*tag= d
XX FT /note= "Flanking sequence"
XX
XX WO9416716-A.
XX
XX 04-AUG-1994.
XX
XX 21-JAN-1994; 94WO-US00888.
XX
XX 21-JAN-1993; 93US-0007115.
XX
XX 19-JAN-1994; 94US-0184009.
XX
XX (VIRO-) VIROGENETICS CORP.
XX
XX Cox WI, Paoletti E, Tartaglia J;
XX
XX WPI; 1994-263767/32.
XX
XX Attenuated recombinant virus used for cancer therapy - comprises
XX DNA encoding cytokine and/or tumour associated antigen
XX
XX Example 16; Fig 20; 232pp; English.
XX
XX The sequences given in Q67865-66 represent expression cassettes
XX containing the vaccinia H6 promoter and the human MAGE-1 gene which
XX encodes human melanoma-associated antigen MZ2-E, in vCP235 and pMAW037,
XX respectively. These sequences were used in the construction of NTVAC-
XX and ALVAC-based recombinant viruses containing the MAGE-1 gene. MAGE-1
XX is expressed in primary melanoma tumour cells, melanoma-derived cell
XX lines and certain tumours of non-melanoma origins but not in normal
XX cells except in testis. A first PCR fragment containing the last 18 bp
XX and the initial 24 nucleotides of the MAGE-1 gene was generated and
XX ligated to a second PCR fragment amplified from plasmid PTZ18RWAGE1
XX which contains the initial 546 bp of the MAGE-1 coding sequence. The
XX terminal sequence of MAGE-1 was amplified and a fusion product was
XX generated containing the H6 promoter and the full length MAGE-1 sequence.
XX This construct may be introduced in to the poxvirus derived plasmids,

```

ALVAC and NYVAC. The resulting viruses may be used in a composition for inducing an antigenic or immunological response, ie. for immunisation against pathogens.

Sequence 1084 BP; 266 A; 256 C; 280 G; 282 T; 0 other;

Query Match 0.6%; Score 17; DB 15; Length 1084;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1652 ggctctggggcaccctgg 1668
|||||

292 ggctctggggcaccctgg 308

RESULT 40

Z08442
Z08442 standard; DNA; 1084 BP.

Z08442;

19-OCT-1999 (first entry)

H6/WAGE-1 expression cassette and flanking regions from pMAW037.

Attenuated recombinant virus; cytokine; tumour associated antigen;
NYVAC recombinant virus; ALVAC recombinant virus; gene therapy; rabies;
cancer; tumour necrosis factor; nuclear phosphoprotein; p53; IL-2; GM-CSF;
interleukin; interferon; IFN-gamma; IL-4; melanoma associated antigen;
carcinoembryonic antigen; immunisation; antigenic; poxvirus; influenza;
immunological response; immunotherapy; vaccine; Newcastle Disease; ss.

Synthetic.

Homo sapiens.

Vaccinia virus.

US5942235-A.

24-AUG-1999.

02-JUN-1995; 95US-0458356.

02-JUN-1995; 95US-0458356.

24-DEC-1981; 81US-0334456.

08-DEC-1982; 82US-0446824.

19-JUN-1984; 84US-0622135.

27-AUG-1987; 87US-0090209.

28-OCT-1987; 87US-0090711.

25-APR-1988; 88US-0186054.

23-AUG-1988; 88US-0234390.

08-MAR-1989; 89US-0320471.

14-FEB-1990; 90US-0478179.

14-JUN-1990; 90US-0537882.

14-JUN-1990; 90US-0537890.

07-JAN-1991; 91US-0638080.

07-MAR-1991; 91US-0668056.

11-JUN-1991; 91US-0713967.

16-DEC-1991; 91US-0805567.

03-MAR-1992; 92US-0847977.

06-MAR-1992; 92US-0847951.

04-MAY-1992; 92US-0881995.

22-JUL-1992; 92US-0918278.

20-JAN-1993; 93US-0007115.

19-JAN-1994; 94US-0184009.

14-APR-1994; 94US-0228926.

13-SEP-1994; 94US-0306259.

(HEAL-) HEALTH RES INC.

Paolotti E;

WPI; 1999-493494/41.

Recombinant poxviruses comprising exogenous DNA encoding antigenic determinants useful in immunotherapy to immunize against cancers and other diseases such as influenza, Newcastle Disease and rabies

Example 16; Fig 20; 163pp; English.

The present invention describes a recombinant poxvirus (I), comprising exogenous DNA encoding an antigenic determinant of a pathogen which is then expressed in vivo in infected host cells after administration to a patient and therefore induces an immunological response. (I) may be used to vaccinate patients against a wide range of diseases and disorders depending on the type of antigen encoded by the exogenous DNA. (I) may be used to vaccinate against diseases such as rabies, influenza and Newcastle Disease. It is particularly useful for immunising against cancers. The poxvirus (I) also provides a means of manipulating lymphocytes and tumour cells for use in cell-based immunotherapeutic modalities for cancer. (I) also have enhanced safety compared to unattenuated viruses (attenuation reduces the virulence of the viruses) and known recombinant poxvirus vaccines. This increased level of safety reduces the possibility of a 'runaway' infection in the host and reduces the chance of transmission from vaccinated to unvaccinated individuals and contamination of the environment. The present sequence represents a H6/WAGE-1 expression cassette and flanking regions from pMAW037 used in the exemplification of the present invention.

Sequence 1084 BP; 266 A; 256 C; 280 G; 282 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 1084;

Best Local Similarity 100.0%; Pred. No. 1.7e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggctctggggcaccctgg 1668

|||||

Db 292 ggctctggggcaccctgg 308

RESULT 41

Q67865

ID Q67865 standard; DNA; 1094 BP.

AC Q67865;

DT 22-MAR-1995 (first entry)

DE H6/WAGE-1 expression cassette from VCP235.

Polymerase chain reaction; primer; amplify; NYVAC; ALVAC; recombinant; human; MAGE-1; melanoma-associated antigen; M22-E; testis; pTZ18RMAGE1; primary melanoma tumour cell; melanoma-derived cell line; tumour; poxvirus; antigenic response; immunological response; pathogen; ss.

OS Synthetic.

FH Key Location/Qualifiers

Promoter 74..200

FT /*tag= a

FT /*note= "vaccinia H6 promoter"

FT CDS 201..1031

FT /*tag= b

FT /*product= MAGE-1

FT misc_feature 1032..1094

FT /*tag= c

FT /*note= "Flanking sequence"

PN WO9416716-A.

PD 04-AUG-1994.

XX 21-JAN-1994; 94WO-US00888.

PF

11-JUN-1999 (first entry)
Human secreted protein gene 43 clone HTADX17.
Human; secreted protein; fusion protein; gene therapy; protein therapy;
diagnosis; tissue; cancer; tumour; neurodegenerative disorder; leukaemia;
developmental abnormality; foetal deficiency; blood; allergy; renal; ds;
immune system; asthma; lymphocytic disease; brain; hepatic; lymphoma;
inflammation; ischaemic shock; Alzheimer's disease; restenosis; AIDS;
cognitive disorder; schizophrenia; prostate; obesity; osteoclast; thymus;
osteoporosis; arthritis; testis; lung; thyroiditis; thyroid; digestion;
endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.
Homo sapiens.
WO9902546-A1.
21-JAN-1999.
07-JUL-1998; 98WO-US13684.
12-SEP-1997; 97US-0058785.
08-JUL-1997; 97US-0051916.
08-JUL-1997; 97US-0051918.
08-JUL-1997; 97US-0051919.
08-JUL-1997; 97US-0051920.
08-JUL-1997; 97US-0051925.
08-JUL-1997; 97US-0051926.
08-JUL-1997; 97US-0051928.
08-JUL-1997; 97US-0051929.
08-JUL-1997; 97US-0051930.
08-JUL-1997; 97US-0051931.
08-JUL-1997; 97US-0051932.
08-JUL-1997; 97US-0052732.
08-JUL-1997; 97US-0052733.
08-JUL-1997; 97US-0052793.
08-JUL-1997; 97US-0052795.
08-JUL-1997; 97US-0052803.
18-AUG-1997; 97US-0055684.
18-AUG-1997; 97US-0055722.
18-AUG-1997; 97US-0055723.
18-AUG-1997; 97US-0055947.
18-AUG-1997; 97US-0055948.
18-AUG-1997; 97US-0055949.
18-AUG-1997; 97US-0055950.
18-AUG-1997; 97US-0055953.
18-AUG-1997; 97US-0055954.
18-AUG-1997; 97US-0055964.
18-AUG-1997; 97US-0055984.
18-AUG-1997; 97US-0056360.
12-SEP-1997; 97US-0058660.
12-SEP-1997; 97US-0058661.
12-SEP-1997; 97US-0058664.
(HUMA-) HUMAN GENOME SCI INC.
Brewer LA, Ebner R, Fischer CL, Kwaw H, Lafleur DW, Li Y, Moore PA;
Olsen HS, Rosen CA, Ruben SM, Shi Y, Soppet DR, Zeng Z;
WPI; 1999-120770/10.
P-PSDB; Y02692.
New isolated human genes and the secreted polypeptides they encode -
useful for diagnosis and treatment of e.g. cancers, neurological
disorders, immune diseases, inflammation or blood disorders
Claim 1; Page 271; 464pp; English.
This sequence represents a nucleic acid molecule which encodes a secreted
human protein. The gene number, and the clone it is derived from, are
detailed in the descriptor line. The gene can be used to generate fusion
proteins by linking to the gene to a human immunoglobulin Fc portion
(e.g. X27302) for increasing the stability of the fused protein as
compared to the human protein only.
The invention relates to 123 novel genes and their fragments (nucleic
acid sequences: X27311-X27449; amino acid sequences Y02650-Y02788) which
are useful for preventing, treating or ameliorating medical conditions
e.g. by protein or gene therapy. Also, pathological conditions can be
diagnosed by determining the amount of the new polypeptides in a sample
or by determining the presence of mutations in the new polynucleotides.
Specific uses are described for each of the 123 polynucleotides, based on
which tissues they are most highly expressed in (see X27311 for described
uses).
Sequence 1140 BP; 280 A; 312 C; 289 G; 254 T; 5 other;
Query Match 0.6%; Score 17; DB 20; Length 1140;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1802 caaagcctggctccagc 1818
DB 930 caaagcctggctccagc 946
|||||
RESULT 44
Q29634
ID Q29634 standard; DNA; 1173 BP.
XX Q29634;
XX
XX 16-MAR-1993 (first entry)
XX Hepatitis C virus HC-J5 3' region.
XX Hepatitis C virus.
XX Non-A non-B hepatitis; NANBH; HCV; detection; diagnosis; screening;
XX PCR; primer; polymerase chain reaction; ss.
XX
XX Hepatitis C virus.
XX EP510952-A.
XX
XX 28-OCT-1992.
XX
XX 23-APR-1992; 92EP-0303625.
XX
XX 26-APR-1991; 91JP-0191376.
XX
XX (IMMO) IMMUNO JAPAN INC.
XX
XX Nakamura T, Okamoto H;
XX
XX WPI; 1992-359137/44.
XX
XX Detection of non-A, non-B hepatitis virus - using new
XX oligo-nucleotide primers with nucleotide sequences corresp. to
XX part. of the viral RNA
XX
XX Disclosure; Page 28; 54pp; English.
XX
XX This sequence represents the 3' region of hepatitis C virus RNA. The
XX original sample was obtained from human and chimpanzee plasma. RNA
XX was isolated from several samples and homology compared, and the
XX respective sequence of about 1900 - 2500 nucleotides of the 5'
XX terminus and 1100 nucleotides of the 3' terminus determined.
XX
XX Sequence 1173 BP; 246 A; 358 C; 310 G; 259 T; 0 other;
Query Match 0.6%; Score 17; DB 13; Length 1173;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 294 cggatctcttcaactg 310
|||||

```

D 910 cggatctcttcaactg 926
RESULT 45
D Q29635 standard; DNA; 1173 BP.
C Q29635;
X 16-MAR-1993 (first entry)
X Hepatitis C virus HC-J6 3' region.
X Non-A non-B hepatitis; NANBH; HCV: detection; diagnosis; screening;
X PCR; primer; polymerase chain reaction; ss.
X Hepatitis C virus.
X EP510952-A.
X 28-OCT-1992.
X 23-APR-1992; 92EP-0303625.
X 26-APR-1991; 91JP-0191376.
X (IMMO) IMMUNO JAPAN INC.
X Nakamura T, Okamoto H;
X WPI; 1992-359137/44.
X Detection of non-A, non-B hepatitis virus - using new
X oligo-nucleotide primers with nucleotide sequences corresp. to
X part. of the viral RNA
X Disclosure; Page 29; 54pp; English.
X This sequence represents the 3' region of hepatitis C virus RNA. The
X original sample was obtained from human and chimpanzee plasma. RNA
X was isolated from several samples and homology compared, and the
X respective sequence of about 1900 - 2500 nucleotides of the 5'
X terminus and 1100 nucleotides of the 3' terminus determined.
X Sequence 1173 BP; 251 A; 362 C; 304 G; 256 T; 0 other;

Query Match 0.6%; Score 17; DB 13; Length 1173;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 294 cggatctcttcaactg 310
D 910 cggatctcttcaactg 926
RESULT 46
D Q43895 standard; cDNA to RNA; 1173 BP.
C Q43895;
X 21-OCT-1993 (first entry);
X NANB hepatitis virus polynucleotide N-1173-3.
X Non-A, non-B; virus; polymerase chain reaction; detection;
X sensitive; specific; HCV; NANBH; ss.
X Non-A, non-B hepatitis virus.
X Key Location/Qualifiers
X CDS 2..1119

Query Match 0.6%; Score 17; DB 13; Length 1173;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 294 cggatctcttcaactg 310
D 910 cggatctcttcaactg 926
RESULT 47
D Q46089 standard; cDNA; 1209 BP.
C Q46089;
X 05-MAY-2000 (first entry)
X cDNA encoding a forkhead activin signal transducer designated FAST2.
X Forkhead activin signal transducer protein; FAST2; activin signalling;
X winged-helix/forkhead domain protein; homeobox gene; goosecoid inducer;
X gsc; transforming growth factor-beta signalling; gsc promoter;
X signal transduction; transcription factor; wound healing; inflammation;
X tumour progression; scarring; arthritis; fibrosis; liver fibrosis;
X kidney fibrosis; ss.
X Mus sp.
X Key Location/Qualifiers
X CDS 4..1209
X /tag= a
X /product= "Forkhead activin signal transducer protein"
X /transl_except= (pos: 1183..1185, aa: Cys)
X WO200004143-A1.
X 27-JAN-2000.
X 19-JUL-1999; 99WO-CA00645.
X

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FT XX
PN XX JP05091884-A.
XX PD 16-APR-1993.
XX PF 10-APR-1991; 91JP-0196175.
XX PR 12-JUN-1990; 90JP-0153401.
XX PR 08-NOV-1990; 90JP-0304405.
XX PA (NAKA/) NAKAMURA T.
XX WPI; 1993-199637/25.
XX P-PSDB; R38285.
XX Antigen related to non-A and non-B hepatitis virus - comprises
XX non-translation region comprising 340 - 341 mols. of nucleotides,
XX non-translation region comprising 1885 - 2551 mols. of
XX nucleotides including region 1,149 and, etc.
XX Claim 9; Page 29; 73pp; Japanese.
XX The sequence is that of NANB hepatitis virus polynucleotide N-1173-3
XX which codes for a non-A, non-B (NANB) hepatitis virus gene HC-OM.
XX The polypeptide it encodes may be used in a system for detecting
XX NANB hepatitis. This method is highly specific and sensitive, and
XX can detect NANB hepatitis virus which could not be detected by
XX conventional methods.
XX Sequence 1173 BP; 247 A; 358 C; 309 G; 259 T; 0 other;

Query Match 0.6%; Score 17; DB 14; Length 1173;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 294 cggatctcttcaactg 310
D 910 cggatctcttcaactg 926
RESULT 47
D Q46089 standard; cDNA; 1209 BP.
C Q46089;
X 05-MAY-2000 (first entry)
X cDNA encoding a forkhead activin signal transducer designated FAST2.
X Forkhead activin signal transducer protein; FAST2; activin signalling;
X winged-helix/forkhead domain protein; homeobox gene; goosecoid inducer;
X gsc; transforming growth factor-beta signalling; gsc promoter;
X signal transduction; transcription factor; wound healing; inflammation;
X tumour progression; scarring; arthritis; fibrosis; liver fibrosis;
X kidney fibrosis; ss.
X Mus sp.
X Key Location/Qualifiers
X CDS 4..1209
X /tag= a
X /product= "Forkhead activin signal transducer protein"
X /transl_except= (pos: 1183..1185, aa: Cys)
X WO200004143-A1.
X 27-JAN-2000.
X 19-JUL-1999; 99WO-CA00645.
X

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17-JUL-1998; 98CA-2237788.
 (HSCR-) HSC RES & DEV LP.
 Wrana JL, Attisano L;
 WPI: 2000-171267/15.
 P-PSDB; Y54601.

New mammalian transcription factor, useful for preventing or treating disorders associated with transforming growth factor beta or activin signaling pathways

Claim 3; Page 43; 76pp; English.

The present sequence encodes a mammalian forkhead activin signal transducer (FAST) protein, designated FAST2. The protein is a winged-helix/forkhead domain protein. The protein is an inducer of the homeobox gene goosecoid (gsc) by transforming growth factor (TGF)-beta or activin signaling. FAST2 binds to a nucleotide sequence in the gsc promoter. The FAST2 protein is useful for modulating signal transduction in a TGF-beta or activin signaling pathway, which involves FAST2 as transcription factor, by modulating the formation of Smad2/Smad4/FAST2 or Smad3/Smad4/FAST2 complex. Inhibition of FAST2 binding to its target DNA site inhibits FAST2 specific TGF-beta signaling, which is associated with wound healing inflammation, and tumour progression. Excessive signaling is associated with scarring, arthritis and fibrosis in numerous diseases, including fibrosis of the liver and kidney.

Sequence 1209 BP; 247 A; 412 C; 315 G; 235 T; 0 other;

Query Match 0.6%; Score 17; DB 21; Length 1209;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1074 cacatggccccagcacc 1090
 636 cacatggccccagcacc 652

RESULT 48
 X87593
 X87593 standard; cDNA; 1338 BP.
 X87593;
 26-OCT-1999 (first entry)
 CLYTA-MAGE-1-His fusion DNA.
 MAGE-1; CLYTA-MAGE-1-His; fusion protein; tumour; melanoma;
 breast cancer; bladder cancer; lung cancer; colon cancer;
 head and squamous cell carcinoma; oesophagus carcinoma; vaccine;
 human; ss.
 Chimeric - Streptococcus pneumoniae.
 Chimeric - Homo sapiens.
 Chimeric - synthetic.
 WO9940188-A2.
 12-AUG-1999;
 02-FEB-1999; 99WO-EP00660.
 06-FEB-1998; 98GB-0002650.
 05-FEB-1998; 98GB-0002543.
 (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
 Cabezón Silva T, Cohen J, Slaoui MM, Vinals Bassols C;

XX WPI: 1999-494293/41.
 DR P-PSDB; Y06592.
 XX
 PT New protein derivatives used in cancer vaccine therapy for treating
 PT a range of cancers including melanomas, carcinomas and cancers of
 PT breast
 XX
 PS Example 9; Page 70-71; 72pp; English.
 XX
 CC This DNA sequence codes for a fusion protein (see Y06592) composed
 CC of the C-terminal portion of the Streptococcus pneumoniae LYTA
 CC protein (CLYTA), the human MAGE-1 tumour-associated antigen and a
 CC hexahistidine tail. A vector designed for recombinant expression
 CC of the fusion protein in Escherichia coli is provided. The CLYTA
 CC moiety provides expression of soluble fusion protein, facilitates
 CC affinity purification of the fusion protein, and also acts as a
 CC T-helper epitope. The invention relates to MAGE proteins fused to
 CC an immunological fusion partner, e.g. CLYTA-MAGE-1-His. These novel
 CC fusion proteins provide vaccines for immunotherapy of melanomas or
 CC other MAGE-associated tumours like breast, bladder, lung and
 CC non-small cell lung cancer, head and squamous cell carcinoma, colon
 CC carcinoma and oesophagus carcinoma.
 XX Sequence 1338 BP; 335 A; 334 C; 378 G; 291 T; 0 other;
 SQ

Query Match 0.6%; Score 17; DB 20; Length 1338;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1652 ggtcctggggcaccctgg 1668
 495 ggtcctggggcaccctgg 511

RESULT 49
 X87591
 ID X87591 standard; cDNA; 1341 BP.
 XX
 AC X87591;
 XX
 DT 26-OCT-1999 (first entry)
 XX
 DE Lipoprotein D-MAGE-1-His fusion DNA.
 XX
 KW MAGE-1; lipoprotein D; LPD-MAGE-1-His; fusion protein; tumour;
 KW melanoma; breast cancer; bladder cancer; lung cancer;
 KW head and squamous cell carcinoma; colon cancer;
 KW oesophagus carcinoma; vaccine; human; ss.
 XX
 OS Chimeric - Haemophilus influenzae.
 OS Chimeric - Homo sapiens.
 OS Chimeric - synthetic.
 XX
 PN WO9940188-A2.
 XX
 PD 12-AUG-1999.
 XX
 PF 02-FEB-1999; 99WO-EP00660.
 XX
 PR 06-FEB-1998; 98GB-0002650.
 PR 05-FEB-1998; 98GB-0002543.
 XX
 PA (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
 XX
 PI Cabezón Silva T, Cohen J, Slaoui MM, Vinals Bassols C;
 XX
 DR WPI: 1999-494293/41.
 DR P-PSDB; Y06590.
 XX
 PT New protein derivatives used in cancer vaccine therapy for treating
 PT a range of cancers including melanomas, carcinomas and cancers of

breast

Example 6; Page 66; 72pp; English.

This DNA sequence codes for a fusion protein (see Y06590) composed of lipidated protein D (LPD) of Haemophilus influenzae B, the human MAGE-1 tumour-associated antigen and a hexahistidine tail. The invention relates to MAGE proteins fused to an immunological fusion partner such as LPD. The LPD moiety provides the fusion protein with additional exogenous T-cell epitopes and also increase expression levels in E. coli. The lipid tail ensures optimal presentation of the antigen to antigen-presenting cells. The affinity tag facilitates purification. The novel fusion proteins provide vaccines for immunotherapy of melanomas or other MAGE-associated tumours like breast, bladder, lung and non-small cell lung cancer, head and squamous cell carcinoma, colon carcinoma and oesophagus carcinoma.

Sequence 1341 BP; 336 A; 327 C; 351 G; 327 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 1341;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1652 ggtcctggggcaccctgg 1668
|||||
498 ggtcctggggcaccctgg 514

RESULT 50
114450/C
Q14450 standard; RNA; 1345 BP.

Q14450;
11-DEC-1991 (first entry)
16S RNA from ATCC 19588 sulphate-reducing bacteria.
SRB; Desulphovibrio; Desulphotomaculum; ribosomal RNA; ss.
Synthetic.

Key Location/Qualifiers
US049489-A.
17-SEP-1991.
17-APR-1989; 89US-0339277.
17-APR-1989; 89US-0339277.
(STAH) STANDARD OIL CO.
(OHIO) OHIO OIL CO.
Aldrich KJ, Brink DE;
WPI; 1991-294983/40.
Assay for sulphate-reducing bacteria - by hybridisation using a labelled oligo-nucleotide probe corresponding to 16S rRNA of the bacteria

Disclosure; Fig 1; 16pp; English.

The sequence is shown folded into secondary structure in the specification. It was compared with other sequences available in the literature to design probes specific for SRB. The probes can be used for the rapid identification and quantification of SRB in a sample, e.g. oil-field prodn. waters, water from water treatment facilities, or samples from the gut of ruminant animals.

CC See also Q13729-Q13733.
XX Sequence 1345 BP; 331 A; 297 C; 435 G; 249 U; 33 other;
SQ

Query Match 0.6%; Score 17; DB 12; Length 1345;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 20 cggcttctcagtttg 36
|||||
Db 1274 CGGCTTCTCAGTTTG 1258

RESULT 51
T27644/C
ID: T27644 standard; cDNA; 1557 BP.

AC T27644;
DT 14-NOV-1996 (first entry)
XX cDNA encoding protein for releasing G1 arrest in an animal cell.
DE G1 arrest; animal cell; human foreskin; cell cycle; ds.
KW Homo sapiens.
OS
XX
FH Key Location/Qualifiers
FT CDS 280..1290
FT /*tag= a
FT Misc-difference 1436
FT /*tag= b
FT /note= "Given in the specification as 5"

PN JP08092288-A.
XX
PD 09-APR-1996.
PF 21-SEP-1994; 94JP-0251537.
XX
PR 21-SEP-1994; 94JP-0251537.
XX
PA (TEIJ) TEIJIN LTD.
XX
XX WPI; 1996-236098/24.
DR P-PSDB; R96248.
XX
PT DNA encoding G1 arrest-releasing protein - useful for the control of the cell cycle
PT the cell cycle
PS Claim 3; Page 9-10; 12pp; Japanese.
XX
CC This sequence encodes a protein which can release G1 arrest of an animal cell. This sequence was isolated from a human foreskin cDNA library. The protein can be used in the control of the cell cycle.
XX
SQ Sequence 1557 BP; 438 A; 349 C; 363 G; 406 T; 1 other;

Query Match 0.6%; Score 17; DB 17; Length 1557;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 708 aatgagccacaccttc 724
|||||
Db 1014 AATGAGCCACACCTTC 998

RESULT 52
T98595/C
ID T98595 standard; DNA; 1591 BP.

T98595;
06-NOV-1998 (first entry)
DNA encoding GTP-binding proteins ERA homolog.
Streptococcus pneumoniae protein; genetic immunisation; antagonist;
immunological response; inoculation; antibody production; inhibitor;
T cell immune response; antimicrobial compound; bacterial adhesion;
extracellular matrix protein; protein-mediated cell invasion; wound;
pathogenesis; ss.
Streptococcus pneumoniae.
Key Location/Qualifiers
CDS complement (592..1086)
/*tag= a
CDS complement (1022..1492)
/*tag= b
W09743303-A1.
20-NOV-1997.
14-MAY-1997; 97WO-US07950.
14-MAY-1996; 96US-0017670.
(SMIK) SMITHKLINE BEECHAM CORP.
(SMIK) SMITHKLINE BEECHAM PLC.
Black MT, Hodgson JE, Knowles DJC, Nicholas RO;
Stodola RK;
WPI: 1998-008793/01.
P-PSDB; W38537, W38538.
Novel Streptococcus pneumoniae proteins and related DNA - useful for
diagnosing anti-microbial agents for treatment of bacterial
infections
Claim 4; Pages 129-130; 483pp; English.
This sequence encodes two Streptococcus pneumoniae proteins (based
on homology with Streptococcus mutans proteins) are GTP-binding
proteins ERA homolog, and represents a DNA sequence of the invention.
The DNA sequences were isolated from Streptococcus pneumoniae strain
0100993 (NCIMB 40794). The Streptococcus pneumoniae proteins of the
invention can be used to identify compounds which interact with and
inhibit or activate the activity of the proteins. Antagonists can be
used to treat diseases caused by S. pneumoniae proteins, through genetic
immunisation. They can also be used to induce an immunological response
in a mammal by inoculation with the S. pneumoniae proteins or delivery
of the encoding nucleic acids in a vector adequate to produce antibody
and/or T cell immune responses to protect the animal from disease. The
proteins can also be used to identify antimicrobial compounds which are
capable of inhibiting their bioactivity. In particular the proteins of
the invention can be used to prevent adhesion of bacteria to mammalian
extracellular matrix proteins on in-dwelling devices or in wounds, to
block protein-mediated mammalian cell invasion, and to block the normal
progression of pathogenesis in infections initiated other than by the
implantation of in-dwelling devices or other surgical techniques.
Sequence 1591 BP; 452 A; 378 C; 313 G; 448 T; 0 other;
Query Match 0.6%; Score 17; DB 19; Length 1591;
Best Local Similarity 100.0%; Pred. No. 1.7e-02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1947 gatttggagaggtttca 1963
|||||
DB 1158 gatttggagaggtttca 1174

Db 435 GATTTCGAAGAGTTTCA 419
RESULT 53
V42963
ID V42963 standard; DNA; 1592 BP.
XX
AC V42963;
XX
DT 09-NOV-1998 (first entry)
XX
DE Streptococcus pneumoniae polypeptide coding region.
XX
KW Polypeptide; ORF; open reading frame; infection; bacterial;
KW streptococcal; bacteremia; diagnosis; prophylaxis; ds.
XX
OS Streptococcus pneumoniae.
XX
FH Key Location/Qualifiers
CDS 573..1001
FT /*tag= a
FT /*note= "polypeptide"
XX
PN W09823631-A1.
XX
PD 04-JUN-1998.
XX
PF 24-NOV-1997; 97WO-US21976.
XX
PR 27-NOV-1996; 96US-0031879.
XX
PA (SMIK) SMITHKLINE BEECHAM CORP.
PA (SMIK) SMITHKLINE BEECHAM PLC.
XX
PI Black MT, Hodgson JE, Knowles DJC, Lonetto MA, Nicholas RO;
PI Reid RH, Zarfos PN;
XX
DR WPI: 1998-322654/28.
DR P-PSDB; W62683.
XX
PT Streptococcus pneumoniae polynucleotides - useful for developing
PT products for diagnosis, prevention and treatment of infections e.g.
PT pneumonia, bacteremia, meningitis or endocarditis
XX
PS Claim 1; Page 56-57; 181pp; English.
XX
CC The sequence is that of a Streptococcal polypeptide coding region.
CC The polypeptide can potentially be used for the diagnosis and
CC prevention of bacterial infections, especially SP infection.
CC It may be used for the treatment of diseases such as otitis media,
CC conjunctivitis, pneumonia, bacteremia, meningitis, sinusitis, pleural
CC empyema, endocarditis or infection of the cerebrospinal fluid.
XX
SQ Sequence 1592 BP; 454 A; 317 C; 376 G; 445 T; 0 other;
Query Match 0.6%; Score 17; DB 19; Length 1592;
Best Local Similarity 100.0%; Pred. No. 1.7e-02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1947 gatttggagaggtttca 1963
|||||
DB 1158 gatttggagaggtttca 1174
RESULT 54
V68059
ID V68059 standard; DNA; 1632 BP.
XX
AC V68059;
XX
DT 02-FEB-1999 (first entry)
XX

Neurodegenerative polypeptide HHPDZ65var coding sequence.
Neurodegenerative polypeptide; HHPDZ65; stroke; pain; epilepsy; therapy;
neurodegenerative disease; ss.
Homo sapiens.
EP875570-A2.
04-NOV-1998.
15-APR-1998; 98EP-0302912.
19-FEB-1998; 98GB-0003566.
01-MAY-1997; 97GB-0008936.
18-DEC-1997; 97EP-0310289.
(SMIK) SMITHKLINE BEECHAM PLC.
Bingham S, Davis J, Doe TR, Harrison DC, Topp S;
WPI; 1998-559436/48.
P-PSDB; W80318.
HHPDZ65 polypeptide(s), their corresponding DNA, antibodies,
agonists and antagonists - are useful in the treatment of stroke,
pain, epilepsy and neurodegenerative diseases
Claim 21; Page 16; 3lpp; English.
This sequence encodes the HHPDZ65var neurodegenerative
polypeptide of the invention. HHPDZ65 is useful for the treatment of
stroke, pain, epilepsy, neurodegenerative diseases and others. The DNAs
and proteins are useful in a method for screening to identify compounds
which stimulate or inhibit the function of the HHPDZ65 proteins. The
polypeptides are useful in a process for diagnosing a disease or a
susceptibility to a disease in a subject related to expression or
activity of the HHPDZ65 polypeptides.
Sequence 1632 BP; 312 A; 529 C; 497 G; 294 T; 0 other;
Query Match 0.6%; Score 17; DB 19; Length 1632;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
2624 cactgctccccaggagg 2640
|||||
1579 cactgctccccaggagg 1595
RESULT 55
25023
225023 standard; cDNA; 1664 BP.
225023;
06-DEC-1999 (first entry)
Murine D6 encoding cDNA SEQ ID NO:4.
D6; G protein-coupled heptahelical receptor; diagnosis; asthma;
respiratory inflammatory disorder; identification; ss.
Mus sp.
WO9947697-A1.
23-SEP-1999.
19-MAR-1999; 98WO-US06075.
20-MAR-1998; 98US-0045583.

(MILL-) MILLENNIUM PHARM INC.
(CRCT-) CRC TECHNOLOGY LTD.
Graham GJ, Benjamin Nibbs RJ, Gonzalo J, Gutierrez-Ramos J;
WPI; 1999-562123/47.
P-PSDB; Y41682.
Identification of D6 G-protein coupled receptor binding compounds and
modulators, useful in treatment of asthma
Claim 2; Fig 2; 152pp; English.
Methods have been developed for identifying a compound, which binds to a
human or murine D6 protein, an allelic variant or a fragment comprises
detecting binding of the test compound to the protein. Also described in
the present invention are: (1) a method for identifying a compound
capable of treating a disorder characterised by aberrant D6 nucleic acid
expression of D6 protein activity; (2) a method for treating a subject
having a disorder characterized by aberrant D6 protein activity or
nucleic acid expression comprising administering to the subject a D6
modulator such that treatment of the subject occurs; and (3) methods for
identifying a compound that modulates the activity of a Human or murine
D6 protein, an allelic variant or a fragment. The methods are useful for
identifying compounds capable of treating disorders, especially a
respiratory inflammatory disorder, characterized by aberrant D6 nucleic
acid expression or D6 protein activity. In particular, the disorder is
asthma. D6 modulators are used to treat asthma. The present sequence
encodes the murine D6 protein.
Sequence 1664 BP; 313 A; 484 C; 436 G; 431 T; 0 other;
Query Match 0.6%; Score 17; DB 20; Length 1664;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
2212 tctgaacacattcagc 2228
|||||
1479 tctgaacacattcagc 1495
RESULT 56
V72116
ID V72116 standard; cDNA; 1668 BP.
AC V72116;
XX
XX 10-MAY-1999 (first entry)
DE Mouse FAST-1 coding region.
XX
KW Transforming growth factor-beta superfamily signalling; modulator; Smad2;
KW TGF-beta; detection; FAST-1; MH2 domain; Smad interaction domain; SID;
KW treatment; developmental; disorder; immunological; cancer; diagnosis; ss.
OS Mus sp.
XX
XX WO9853830-A1.
XX
PD 03-DEC-1998.
XX
PF 28-MAY-1998; 98WO-US10983.
XX
PR 28-MAY-1997; 97US-0047991.
XX
PA (HARD) HARVARD COLLEGE.
XX
PI Chen X, Whitman M;
XX
DR WPI; 1999-059773/05.
DR P-PSDB; W90249.

Modulating TGF-beta superfamily signalling - comprises use of compounds identified in assays with Smad2, FAST-1 and Smad3, used to develop products for treating, e.g. developmental disorders

Example XII; Page 70; 107pp; English.

This sequence encodes a mouse FAST-1 protein which is used in a method to detect a compound capable of modulating transforming growth factor-beta (TGF-beta) superfamily signalling. The invention describes a complex which forms between FAST-1 and Smad2 and this complex is specifically induced by signals generated by a TGF-beta superfamily member. A domain of FAST-1 directly interacts with Smad2 and this interaction is mediated by specific domains of the two interacting molecules, namely, the MH2 domain of Smad2 and the Smad interaction domain (SID) of FAST-1. The methods and compounds described are useful for the detection and treatment of conditions involving abnormal TGF-beta superfamily signalling. They can be used to treat e.g. developmental disorders, immunological disorders and cancer. The products can also be used for detection and diagnosis.

Sequence 1668 BP; 368 A; 517 C; 432 G; 351 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 1668;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 1074 cacatggccccagcattc 1090
|||||
5 621 cacatggccccagcattc 637

RESULT 57
V69719 standard; cDNA; 1691 BP.

V69719;
01-MAR-1999 (first entry)
Tumour rejection antigen precursor MAGE-A1 cDNA.
MAGE-A1; human; tumour rejection antigen precursor; TRAP;
therapy; diagnosis; ss.
Homo sapiens.
Key Location/Qualifiers
CDS 204..1133
/*tag= a
WO9849184-A1.
05-NOV-1998.
24-APR-1998; 98WO-US08493.
25-APR-1997; 97US-0845528.
(LUDW-) LUDWIG INST CANCER RES.
Boon-Falleur T, De Smet C, Lucas S;
WPI; 1999-024041/02.
P-PSDB; W81548.
Tumour rejection antigen precursors - used for determining presence of cytolytic T cells specific for complexes of a human leukocyte antigen
Disclosure; Page 46-47; 84pp; English.

This nucleotide sequence comprises human tumour rejection antigen precursor (TRAP) MAGE-A1 cDNA, which encodes a 309-amino acid polypeptide (see W81548). MAGE-A1 cDNA shows homology to novel human MAGE-C1 cDNA (see V69720), especially in exons 2 and 3. The open reading frame of MAGE-C1, however, is about 2 kb longer than that of MAGE-A1, most of the difference being accounted for by a large repetitive sequence. MAGE-C1 (see W81546) is a novel member of the MAGE family that may be recognised by cytotoxic T cells, leading to lysis of the tumour cells which express it. It is expressed in a variety of tumours and in normal testis cells, but not by other normal cells. The invention provides MAGE-C1 and MAGE-C2 nucleic acids and polypeptides, useful e.g. in a claimed method for determining the presence of cytolytic T cells specific for complexes of a human leukocyte antigen (HLA).

Sequence 1691 BP; 410 A; 389 C; 465 G; 427 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 1691;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1652 ggtcctggggcaccctgg 1668
|||||
317 ggtcctggggcaccctgg 333

RESULT 58
252876
ID 252876 standard; cDNA; 1702 BP.
XX 252876;
XX 14-MAR-2000 (first entry)
XX Human prostate tumor cDNA library derived EST fragment #19.
XX Pancreas; tumor; EST; expressed sequence tag; human; cytostatic;
XX treatment; ds.
XX Homo sapiens.
XX DE19820190-A1.
XX 04-NOV-1999.
XX 28-APR-1998; 98DE-1020190.
XX 28-APR-1998; 98DE-1020190.
XX (META-) METAGEN GES GENOMFORSCHUNG MBH.
XX Rosenthal A, Specht T, Hinzmann B, Schmitt A, Pilarsky C, Dahl E;
XX WPI; 1999-621386/54.
XX P-PSDB; Y73868, Y73869, Y73870.
XX New human nucleic acid sequences from pancreatic tumors, and related
XX proteins
XX Claim 2; Page 197-198; 502pp; German.
XX This invention describes novel polypeptides and their encoding nucleic
XX acids derived from human pancreatic tumor tissue which have cytostatic
XX activity. The sequences are also useful in producing pharmaceutical
XX compositions for treatment of pancreatic tumors. 252858-253014 represent
XX expressed sequence tag (EST) fragments derived from a human pancreatic
XX tumor cDNA library and which encode the proteins represented in
XX Y73814-Y74252.
XX Sequence 1702 BP; 371 A; 517 C; 476 G; 338 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 1702;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DY 1341 agggagtgccagagga 1357
 |||||
 1112 agggagtgccagagga 1128

RESULT 59
 V38385
 V38385 standard; cDNA; 1724 BP.
 AC V38385;
 TT 24-NOV-1998 (first entry)
 X Beta(1 -> 4)-N-acetylglucosaminyl-transferase (Gnt-IV)b encoding cDNA.
 DE Beta(1 -> 4)-N-acetylglucosaminyl-transferase; Gnt-IV; bovine; human;
 KW enzyme; sugar chain subunit; branched oligosaccharide; polysaccharide;
 XW drug; reagent; food; biopolymer; glycoprotein; erythropoietin; ss.
 X Homo sapiens.
 S
 X Key Location/Qualifiers
 T CDS 43..1689
 T /*tag= a
 T /product= "Gnt-IVb enzyme"
 X W09826053-A1.
 PN 18-JUN-1998.
 FD 10-DEC-1997; 97MO-JP04546.
 X 18-JUN-1997; 97JP-0161462.
 X 12-DEC-1996; 96JP-0332411.
 X (KIRI) KIRIN BEER KK.
 X Minowa M, Oguri S, Takeuchi M, Taniguchi N, Yoshida A;
 WPI; 1998-348516/30.
 X P-PSDB; W63559.
 X Recombinant beta(1-4)-N-acetylglucosaminyl-transferase - allows
 production of difficultly accessible branched poly:saccharides for
 food and drug use
 X Claim 16; Pages 70-74; 112pp; Japanese.
 X This cDNA encodes a human beta(1 -> 4)-N-acetylglucosaminyl-transferase
 (Gnt-IV)b enzyme. The invention provides bovine and human Gnt-IV enzymes
 that can be used for converting sugar chain subunits having one structure
 to another structure. Vectors containing the DNA sequences encoding these
 enzymes can be used to transform host cells for the production of the
 Gnt-IV enzymes. The enzymes are useful in the production of branched
 oligosaccharides and polysaccharides which are difficult of access by
 other methods. They are also useful in the production of drugs, reagents
 and foods and in modifying the properties of biopolymers containing sugar
 chains. The enzyme may also be used for the preparation of glycoproteins
 such as erythropoietin.
 X Sequence 1724 BP; 353 A; 553 C; 501 G; 317 T; 0 other;

Query Match 0.6%; Score 17; DB 19; Length 1724;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DY 1341 agggagtgccagagga 1357
 |||||
 1112 agggagtgccagagga 1128

DB 1565 agggagtgccagagga 1581

RESULT 60
 X85940
 ID X85940 standard; DNA; 1816 BP.
 XX AC X85940;
 XX
 DT 13-SEP-1999 (first entry)
 DE DNA encoding human cell division regulator (HCDR) 1.
 KW Human cell division regulator; HCDR: interphase; inflammation;
 KW cell proliferation; apoptosis; neurodegenerative;
 KW neurodegenerative disease; aplastic anaemia; ischaemic injury;
 KW liver damage; viral infection; hepatitis B; hepatitis C; ss.
 XX Homo sapiens.
 OS
 XX US5928899-A.
 PW
 XX 27-JUL-1999.
 PD
 XX
 PF 01-OCT-1998; 98US-0165234.
 XX
 PR 15-OCT-1997; 97US-0951148.
 PR 01-OCT-1998; 98US-0165234.
 XX
 PA (INCY-) INCYTE PHARM INC.
 XX
 PI Bandman O, Corley NC, Hillman JL, Lal P, Shah P;
 DR WPI; 1999-429499/36.
 DR P-PSDB; Y23782.
 XX
 PT Cell division regulators active in interphase
 XX
 PS Example 1; Fig 1A-E; 59pp; English.
 XX
 CC The present sequence encodes human cell division regulator (HCDR) 1.
 CC HCDR proteins are active in interphase, and are used for the
 CC treatment or prevention of inflammation and disorders associated with
 CC cell proliferation and apoptosis. HCDR may be administered to a
 CC patient having a disorder associated with an increase in apoptosis,
 CC such a disorder may be e.g. neurodegenerative, a neurodegenerative
 CC disease, aplastic anaemia, an ischaemic injury, liver damage, or a
 CC viral infection such as hepatitis B or C.
 XX
 SQ Sequence 1816 BP; 494 A; 473 C; 494 G; 355 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 1816;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DY 804 ggaactctcttggtgct 820
 |||||
 717 ggaactctcttggtgct 733

DB 717 ggaactctcttggtgct 733

RESULT 61
 X01577
 ID X01577 standard; DNA; 1818 BP.
 XX AC X01577;
 XX
 DT 04-MAY-1999 (first entry)
 XX
 DE Human HCDR-1 coding sequence.
 XX
 KW Human; HCDR-1; HCDR-2; HCDR-3; human cell division regulator; apoptosis;
 KW inflammation; cell proliferation disorder; adenocarcinoma; AIDS; ss.

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XX Homo sapiens.
XX US5871973-A.
XX 16-FEB-1999.
XX 15-OCT-1997; 97US-09511148.
XX 15-OCT-1997; 97US-09511148.
XX (INCY-) INCYTE PHARM INC.
XX Bandman O, Corley NC, Hillman JL, Lal P, Shah P;
XX WPI; 1999-166646/14.
XX P-PSDB; W73971.
XX New polynucleotides encoding human cell division regulators (HCDR)
XX - useful for diagnosing, preventing and treating inflammation and
XX disorders associated with cell proliferation and apoptosis
XX Claim 4; Fig 1; 59pp; English.
XX This sequence encodes the human cell division regulator-1 (HCDR-1)
XX protein of the invention. Polynucleotides complementary to the HCDR-1
XX coding sequence can be used as probes to detect the DNA in a sample. The
XX polynucleotide sequences encoding HCDR may be used to prevent/treat
XX inflammation and disorders associated with cell proliferation and
XX apoptosis and in assays that detect activation of cancers.
XX Polynucleotides encoding HCDR may be used for the diagnosis of conditions
XX associated with expression of HCDR, including disorders associated with
XX cell proliferation/apoptosis e.g. adenocarcinoma and AIDS. The
XX polynucleotides may also be used in Southern or Northern analysis, dot
XX blot, or other membrane based technologies; in PCR technologies; or in
XX dipstick, pin, or ELISA assays or microarrays utilising fluids or tissues
XX from patient biopsies to detect altered HCDR expression.
XX Sequence 1818 BP; 494 A; 473 C; 494 G; 355 T; 2 other;

Query Match 0.6%; Score 17; DB 20; Length 1818;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

XX 804 ggaactcttctgtgct 820
XX |||||||
XX 719 ggaactcttctgtgct 735

RESULT 62
XX A15550/c
XX A15550 standard; cDNA to mRNA; 1906 BP.
XX A15550;
XX 31-JUL-2000 (first entry)
XX Human TRAF four associated factor TRAF2 coding sequence.
XX TRAF2; TRAF four associated factor 2; tumour formation; breast cancer;
XX TRAF4; TNF receptor associated factor; tumour diagnosis; ds.
XX Homo sapiens.
XX Key Location/Qualifiers
XX CDS 13..1230
XX /*tag= a
XX /product= TRAF2
XX CA2245340-A1.
XX 19-FEB-2000.

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XX 19-AUG-1998; 98CA-2245340.
XX 19-AUG-1998; 98CA-2245340.
XX (MEDI-) MEDICAL & BIOLOGICAL LAB CO LTD.
XX Yano M, Toji S, Tamai K;
XX WPI; 2000-351124/31.
XX P-PSDB; Y94209.
XX Novel tumour necrosis factor receptor associated factor 4 associated
XX factors useful for developing cancer screens, and treating tumours -
XX Claim 4; Page 47-50; 68pp; English.
XX The present sequence is the coding sequence of human TRAF four
XX associated factor TRAF2. The gene was discovered by screening a human
XX placenta cDNA library using a two-hybrid system. The protein associates
XX with the TRAF domain located at the carboxyl-terminal of TNF
XX receptor associated factor 4 (TRAF4), which is believed to be an
XX oncoprotein. Antibodies that bind to TNF four associated factors (TRAFs)
XX may be used to treat or diagnose tumours (e.g. breast cancer) when
XX labelled with an isotope or an appropriate drug, precursor or enzyme.
XX Antagonists, agonists, and antisense sequences of TRAFs may be used to
XX treat cancers. TRAF proteins, antibodies that recognise them and DNAs
XX for them may be useful as tools for cancer research.
XX Sequence 1906 BP; 655 A; 320 C; 372 G; 559 T; 0 other;

Query Match 0.6%; Score 17; DB 21; Length 1906;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

XX 2796 gtttttaagaagtctt 2812
XX |||||||
XX 582 GTTTTAAAGAGACTCT 566

RESULT 63
XX T09084/c
XX ID T09084 standard; cDNA to mRNA; 2160 BP.
XX AC T09084;
XX XX 09-MAY-1996 (first entry)
XX DT Nitrite reductase gene.
XX DE Nitrite reductase gene.
XX XX nitrite reductase; transgenic crop; transgenic tree; detoxification;
XX KW atmospheric pollutant; ds.
XX KW Populus nigra L. var italica.
XX OS
XX XX Key Location/Qualifiers
XX FH CDS 40..1806
XX FT /*tag= a
XX FT /product= nitrite_reductase
XX FT primer_bind 367..390
XX FT /*tag= b
XX FT primer_bind 838..860
XX FT /*tag= c
XX FT polyA_signal 2114..2119
XX FT /*tag= d
XX FT misc_feature 2146..2160
XX FT /*tag= e
XX FT /label= polyA_site
XX XX JF07236486-A.
XX PN 12-SEP-1995.
XX PD

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XX 02-MAR-1994; 94JP-0032359.
XX 02-MAR-1994; 94JP-0032359.
XX (TOYT ) TOYOTA JIDOSHA KK.
XX WPI: 1995-347454/45.
XX P-PSDB; R87973.
XX Nitrite reductase gene from Populus nigra - useful for breeding
XX trees to remove nitrogen di:oxide from the atmosphere or for crops
XX which produce fewer carcinogenic nitrosamine(s)
XX Claim 1; Page 5-8; 8pp; Japanese.
XX The gene encodes a nitrite reductase gene isolated from Populus nigra.
XX The gene is useful in generation of street trees with a high power for
XX clarifying NO2, an atmospheric pollutant. The gene can also be used in
XX transgenic plants, esp. crops, bred to contain less nitrosamines (a
XX carcinogenic substance present in food).
XX Sequence 2160 BP; 660 A; 388 C; 570 G; 542 T; 0 other;
XX
XX Query Match 0.6%; Score 17; DB 16; Length 2160;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1711 gcttgccaagtattcttg 1727
XX |||||
XX 495 GCTTGCCAAGTATCTTG 479
XX
XX RESULT 64
XX 50582/c
XX 23-MAY-2000 (first entry)
XX Human epidermal protein-6 cDNA.
XX Human epidermal protein-6; HEPI; epithelial disorder; scabies;
XX dyshidrotic eczema; cell proliferative disorder; actinic keratosis;
XX arteriosclerosis; autoimmune deficiency disorder; inflammatory disorder;
XX acquired immune deficiency syndrome; AIDS; Addison's disease; antiHIV;
XX dermatological; anicarteriosclerotic; antiinflammatory;
XX immunosuppressive; ss.
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 47..1636
XX /*tag= a
XX /product= "Human epidermal protein-6"
XX sig_peptide 1..142
XX /*tag= b
XX mat_peptide 148..1636
XX /*tag= c
XX /product= "Mature human epidermal protein-6"
XX
XX WO200006727-A2.
XX 10-FEB-2000.
XX 27-JUL-1999; 99WO-US17107.
XX 28-JUL-1998; 98US-0155203.
XX 07-DEC-1998; 98US-0155254.
XX (INCY-) INCYTE PHARM INC.

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XX Tang YT, Lal P, Corley NC, Guegler KJ, Patterson C, Baughn MR;
XX Yue H;
XX WPI: 2000-195295/17.
XX P-PSDB; Y44989.
XX New human epidermal proteins (HEPI-1) to (HEPI-6) useful for the
XX diagnosis, treatment and prevention of epithelial, cell proliferative,
XX and autoimmune inflammatory disorders
XX Claim 7; Page 78-79; 82pp; English.
XX The present cDNA sequence encodes human epidermal protein-6 (HEPI). The
XX cDNA clone is derived from PTHYNO03 library which was constructed using
XX RNA isolated from left parathyroid tissue of a 69-year-old caucasian
XX female during a partial parathyroidectomy. Recombinant vectors
XX comprising HEPI cDNA are introduced into host cells for protein
XX expression. The HEPI proteins are useful for the treatment of epithelial
XX disorders, including dyshidrotic eczema and scabies, cell proliferative
XX disorders including actinic keratosis and arteriosclerosis, and
XX autoimmune/inflammatory disorders like acquired immune deficiency
XX syndrome (AIDS) and Addison's disease. Pharmaceutical compositions
XX comprising HEPI proteins are useful for treating disorders associated
XX with altered HEPI expression.
XX Sequence 2284 BP; 452 A; 775 C; 534 G; 523 T; 0 other;
XX
XX Query Match 0.6%; Score 17; DB 21; Length 2284;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1881 caggaagggtgagat 1897
XX |||||
XX 2262 CAGGAAGGCTGAGAT 2246
XX
XX RESULT 65
XX X84103
XX ID X84103 standard; DNA; 2418 BP.
XX AC X84103;
XX DT 08-SEP-1999 (first entry)
XX DE E antigen precursor gene.
XX KW Tumour rejection antigen; vaccine; cancer; E antigen precursor gene; ss.
XX OS Homo sapiens.
XX PN US925729-A.
XX PD 20-JUL-1999.
XX PF 02-MAY-1994; 94US-0142368.
XX PR 02-MAY-1994; 94US-0142368.
XX PR 23-MAY-1991; 91US-0705702.
XX PR 09-JUL-1991; 91US-0728838.
XX PR 23-SEP-1991; 91US-0764365.
XX PR 12-DEC-1991; 91US-0807043.
XX (LUDW-); LUDWIG INST CANCER RES.
XX PA Boon T; Chomez P, De Plaen E, Lurquin C, Traversari C;
XX Van Den Eynde B, Van Der Bruggen P, Van Pel A;
XX WPI: 1999-418294/35.
XX New tumour rejection antigen is useful as a vaccine against
XX cancerous diseases

```

XX Example 20; Column 15-18; 58pp; English.

CC This sequence represents the E antigen precursor gene.

CC The invention relates to a tumour rejection antigen sequence that is

CC useful as a tumour rejection antigen for vaccination against cancerous

CC conditions.

XX Sequence 2418 BP; 562 A; 582 C; 675 G; 599 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 2418;

Best Local Similarity 100.0%; Pred. No. 1.7e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctggggcacctgg 1668

Db 738 ggtcctggggcacctgg 754

RESULT 66

Q22351

Q32351 standard; DNA; 2419 BP.

AC Q32351;

XX 22-APR-1993 (first entry)

XX Antigen E gene.

XX Stable; antigen: E; D; F; A; human; melanoma; cell line; MZ2-MEL;

XX cytolytic T cell; MEL3.1; open reading frame; homology; MAGE;

XX melanoma antigen; ss.

XX Homo sapiens.

XX WO9220356-A.

XX 26-NOV-1992.

XX 22-MAY-1992; 92WO-US04354.1.

XX 23-MAY-1991; 91US-0705702.

XX 03-JUL-1991; 91US-0728838.

XX 23-SEP-1991; 91US-0764364.

XX 12-DEC-1991; 91US-0807043.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Boon T, Chomez P, De Plaen E, Lurquin C, Traversari C;

XX Van Den Eynde B, Van Der Bruggen P, Van Pel A;

XX WPI; 1992-415460/50.

XX Nucleic acid mol. encoding a human tumour rejection antigen

XX precursor - useful as an immunostimulant in a vaccine for

XX treating and preventing cancers, also useful in diagnosis

XX Disclosure; Page 69-70; 142pp; English.

XX This sequence encodes the stable antigen E. This antigen is expressed

XX along with antigens "D, F and A" by the human melanoma cell line MZ2-

XX MEL. These antigens are all recognised by cytolytic T cells. A

XX subline of MZ2-MEL is MEL3.1 which only expresses antigen E. This

XX cell line was chosen as a source for the isolation of this sequence.

XX This sequence was found to contain three exons. The open reading frame

XX for antigen E was contained within the first two exons. An ATG is

XX located at position 66 of exon 3 and is followed by an 828 base pair

XX reading frame. The three exons contain 65, 73 and 1551 base pairs.

XX During the isolation of this sequence two different but closely related

XX cDNAs were also identified. These cDNAs, when tested, did not

XX transfer expression of antigen E, but they did show substantial

XX homology to the antigen E cDNA sequence. These new cDNAs represent a

CC new family of genes referred to as melanoma antigens (MAGE) (see also

CC Q32352-69).

XX Sequence 2419 BP; 562 A; 581 C; 677 G; 599 T; 0 other;

Query Match 0.6%; Score 17; DB 13; Length 2419;

Best Local Similarity 100.0%; Pred. No. 1.7e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctggggcacctgg 1668

Db 739 ggtcctggggcacctgg 755

RESULT 67

Q72476

ID Q72476 standard; DNA; 2419 BP.

XX Q72476;

XX 21-JUN-1995 (first entry)

XX Tumour rejection antigen E encoding DNA.

XX Tumour rejection antigen E; melanoma antigen-3; MAGE-3;

XX cancer; cytolytic T cells; antigen D; human leucocyte antigen; ss.

XX Homo sapiens.

XX WO9423031-A.

XX 13-OCT-1994.

XX 17-MAR-1994; 94WO-US02877.

XX 26-MAR-1993; 93US-0037230.

XX (LUDW-) LUDWIG INST CANCER RES.

XX Boon-falleur T, Gaugler B, Van Den Eynde B, Van Der Bruggen P;

XX WPI; 1994-333192/41.

XX New tumour rejection antigen precursor MAGE3 - useful in

XX treatment and diagnosis of cancer

XX Disclosure; Page 58; 105pp; English.

XX Q72476 encodes tumour rejection antigen E, another sequence

XX Q72470 encodes melanoma antigen-3 (MAGE-3) a tumour rejection

XX antigen precursor. Melanomas characterised by the expression of

XX MAGE-3 can be detected, or monitored, by contacting a test sample

XX with an agent that can recognise MAGE-3. The melanoma can be treated

XX by the administration of cytolytic T cells specific for the complex of

XX antigen D (the mature rejection antigen derived from MAGE-3) and a human

XX leucocyte antigen (esp. HLA-A1).

XX Sequence 2419 BP; 562 A; 581 C; 677 G; 599 T; 0 other;

Query Match 0.6%; Score 17; DB 15; Length 2419;

Best Local Similarity 100.0%; Pred. No. 1.7e+02;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctggggcacctgg 1668

Db 739 ggtcctggggcacctgg 755

RESULT 68

ID T05086

ID T05086 standard; DNA; 2419 BP.

```

XX T05086;
AC
XX
XX
XX 26-FEB-1996 (first entry)
XX
XX MZ2-MEL antigen E precursor gene.
XX
XX Melanoma; MZ2-MEL; tumour rejection antigen; cancer; diagnosis; ss.
XX
XX Homo sapiens.
XX
XX W09523874-A1.
XX
XX 08-SEP-1995.
XX
XX 23-FEB-1995; 95WO-US02203.
XX
XX 30-NOV-1994; 94US-0345774.
XX
XX 01-MAR-1994; 94US-0204727.
XX
XX 10-MAR-1994; 94US-0209172.
XX
XX 01-SEP-1994; 94US-0299849.
XX
XX (LUDW-) LUDWIG INST CANCER RES.
XX
XX Boon-Falleur T, Brasseur F, Chomez P, De Plaen E;
XX De Smet C, Gaugler B, Lethe B, Marchand M, Patard J;
XX Szikora J, Van Den Eynde B, Van Derbruggen P, Weynants P;
XX WPI; 1995-320586/41.
XX
XX Determn. of cancerous condition(s) - using a nucleic acid as a
XX primer to determine expression of a MAGE tumour rejection antigen
XX precursor
XX
XX Example 20; Page 69-70; 121pp; English.
XX
XX A gene sequence (T05086) hybridizes with a 2.4 kb fragment from
XX human melanoma cell line MZ2-MEL but not with E- antigen loss
XX variants of MZ2-MEL. This E precursor antigen gene sequence was
XX obtd. from a cosmid derived from DNA of the E+ subclone MZ2-MEL 43.
XX
XX Sequence 2419 BP; 560 A; 581 C; 677 G; 601 T; 0 other;
XX
XX
XX Query Match 0.6%; Score 17; DB 16; Length 2419;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX Y 1652 ggtcctgggacccctgg 1668
XX |||||
XX Db 739 ggtcctgggacccctgg 755
XX
XX RESULT 69
XX X84112
XX Db X84112 standard; DNA; 2419 BP.
XX
XX AC X84112;
XX
XX 08-SEP-1999 (first entry)
XX
XX Antigen E coding sequence:
XX
XX Tumour rejection antigen; vaccine; cancer; antigen E; ss.
XX
XX Homo sapiens.
XX
XX US5925729-A.
XX
XX 20-JUL-1999.
XX
XX 02-MAY-1994; 94US-0142368.
XX

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PR 02-MAY-1994; 94US-0142368.
PR 23-MAY-1991; 91US-0705702.
PR 09-JUL-1991; 91US-0728838.
PR 23-SEP-1991; 91US-0764365.
PR 12-DEC-1991; 91US-0807043.
XX
XX (LUDW-) LUDWIG INST CANCER RES.
XX
XX Boon T, Chomez P, De Plaen E, Lurquin C, Traversari C;
XX Van Den Eynde B, Van Der Bruggen P, Van Pel A;
XX WPI; 1999-418294/35.
XX
XX New tumour rejection antigen is useful as a vaccine against
XX cancerous diseases
XX
XX Disclosure; Column 37-40; 58pp; English.
XX
XX This sequence represents the antigen E coding sequence.
XX The invention relates to a tumour rejection antigen sequence that is
XX useful as a tumour rejection antigen for vaccination against cancerous
XX conditions.
XX
XX Sequence 2419 BP; 562 A; 581 C; 677 G; 599 T; 0 other;
XX
XX
XX Query Match 0.6%; Score 17; DB 20; Length 2419;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1652 ggtcctgggacccctgg 1668
XX |||||
XX Db 739 ggtcctgggacccctgg 755
XX
XX RESULT 70
XX Q72472
XX ID Q72472 standard; DNA; 2420 BP.
XX
XX AC Q72472;
XX
XX DT 21-JUN-1995 (first entry)
XX
XX Tumour rejection antigen E precursor gene DNA.
XX
XX Tumour antigen rejection precursor E; melanoma antigen-3; MAGE-3;
XX cancer; cytolytic T cells; antigen D; human leucocyte antigen; ss.
XX
XX OS Homo sapiens.
XX
XX PN W09423031-A.
XX
XX PD 13-OCT-1994.
XX
XX PF 17-MAR-1994; 94WO-US02877.
XX
XX PR 26-MAR-1993; 93US-0037230.
XX
XX (LUDW-) LUDWIG INST CANCER RES.
XX
XX Boon-falleur T, Gaugler B, Van Den Eynde B, Van Der Bruggen P;
XX WPI; 1994-333192/41.
XX
XX New tumour rejection antigen precursor MAGE3 - useful in
XX treatment and diagnosis of cancer
XX
XX Example 20; Page 28; 105pp; English.
XX
XX Q72472 is the tumour rejection antigen E precursor gene, another
XX gene Q72470 encodes melanoma antigen-3 (MAGE-3) also a tumour rejection
XX antigen precursor. Melanomas characterised by the expression of MAGE-3
XX can be detected, or monitored, by contacting a test sample with an

```

agent that can recognise MAGE-3. The melanoma can be treated by the administration of cytolytic T cells specific for the complex of antigen D (the mature rejection antigen derived from MAGE-3) and a human leucocyte antigen (esp. HLA-A1).

Sequence 2420 BP; 562 A; 582 C; 677 G; 599 T; 0 other;

Query Match 0.6%; Score 17; DB 15; Length 2420;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1652 ggtcctgggacacctgg 1668
|||||

739 ggtcctgggacacctgg 755

RESULT 71

Q85435

Q85435 standard; DNA; 2420 BP.

Q85435;

09-OCT-1995 (first entry)

Human melanoma antigen MAGE-1.

Human melanoma antigen; MAGE-1; vaccines; MAGE associated tumours;
HLA-restricted cytotoxic T-lymphocyte activity; ss.

Homo sapiens.

Key Location/Qualifiers
CDS 626..1555
/*tag= a

W09504542-A.

16-FEB-1995.

02-AUG-1994; 94WO-US08721.

06-AUG-1993; 93US-0103623.

(CYTE-) CYTEL CORP.

Fikes JD, Livingston BD, Sette AD, Sidney JC;

WPI; 1995-090681/12.

P-PSDB; R70909.

Human melanoma antigen, MAGE-1, peptide(s) - useful for
stimulating immune response against melanoma

Example 1; Fig 1; 59pp; English.

Q85435 encodes R70909 human melanoma antigen MAGE-1, it was used to produce the C-terminal MAGE-1 peptides described in R70915 to R70969. These peptides are useful for defining epitopes that engender a HLA-restricted cytotoxic lymphocyte activity against MAGE-1 antigens. Compsns. containing these peptides can be administered, as a vaccine to patients susceptible to MAGE associated tumours, e.g. melanomas.

Sequence 2420 BP; 562 A; 582 C; 677 G; 599 T; 0 other;

Query Match

Best Local Similarity 0.6%; Score 17; DB 16; Length 2420;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1652 ggtcctgggacacctgg 1668
|||||

Db 739 ggtcctgggacacctgg 755
RESULT 72
V59595
ID V59595 standard; DNA; 2503 BP.
XX V59595;
AC V59595;
XX
DT 06-JAN-1999 (first entry)
XX
DE Human secreted protein gene 85 clone HSDPV29.

XX
KW Human; secreted protein; fusion protein; gene therapy; protein therapy;
diagnosis; tissue; cancer; tumour; neurodegenerative disorder; leukaemia;
developmental abnormality; foetal deficiency; blood; allergy; renal; ds;
immune system; asthma; lymphocytic disease; brain; hepatic; lymphoma;
inflammation; ischaemic shock; Alzheimer's disease; restenosis; AIDS;
cognitive disorder; schizophrenia; prostate; obesity; osteoclast; thymus;
osteoporosis; arthritis; testis; lung; thyroiditis; thyroid; digestion;
endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.

Homo sapiens.

W09839448-A2.

11-SEP-1998.

06-MAR-1998; 98WO-US04493.

02-OCT-1997; 97US-0061060.

07-MAR-1997; 97US-0038621.

07-MAR-1997; 97US-0040161.

07-MAR-1997; 97US-0040162.

07-MAR-1997; 97US-0040163.

07-MAR-1997; 97US-0040333.

07-MAR-1997; 97US-0040334.

07-MAR-1997; 97US-0040336.

07-MAR-1997; 97US-0040626.

11-APR-1997; 97US-0043311.

11-APR-1997; 97US-0043312.

11-APR-1997; 97US-0043313.

11-APR-1997; 97US-0043314.

11-APR-1997; 97US-0043568.

11-APR-1997; 97US-0043569.

11-APR-1997; 97US-0043576.

11-APR-1997; 97US-0043578.

11-APR-1997; 97US-0043580.

11-APR-1997; 97US-0043669.

11-APR-1997; 97US-0043670.

11-APR-1997; 97US-0043671.

11-APR-1997; 97US-0043672.

11-APR-1997; 97US-0043674.

23-MAY-1997; 97US-0047492.

23-MAY-1997; 97US-0047500.

23-MAY-1997; 97US-0047501.

23-MAY-1997; 97US-0047502.

23-MAY-1997; 97US-0047503.

23-MAY-1997; 97US-0047581.

23-MAY-1997; 97US-0047582.

23-MAY-1997; 97US-0047583.

23-MAY-1997; 97US-0047584.

23-MAY-1997; 97US-0047585.

23-MAY-1997; 97US-0047586.

23-MAY-1997; 97US-0047587.

23-MAY-1997; 97US-0047588.

23-MAY-1997; 97US-0047589.

23-MAY-1997; 97US-0047590.

23-MAY-1997; 97US-0047592.

23-MAY-1997; 97US-0047593.

23-MAY-1997; 97US-0047594.

23-MAY-1997; 97US-0047595.

23-MAY-1997; 97US-0047596.

23-MAY-1997; 97US-0047597.
 23-MAY-1997; 97US-0047598.
 23-MAY-1997; 97US-0047599.
 23-MAY-1997; 97US-0047600.
 23-MAY-1997; 97US-0047601.
 23-MAY-1997; 97US-0047612.
 23-MAY-1997; 97US-0047613.
 23-MAY-1997; 97US-0047614.
 23-MAY-1997; 97US-0047615.
 23-MAY-1997; 97US-0047617.
 23-MAY-1997; 97US-0047618.
 23-MAY-1997; 97US-0047632.
 23-MAY-1997; 97US-0047633.
 06-JUN-1997; 97US-0048964.
 06-JUN-1997; 97US-0048974.
 13-JUN-1997; 97US-0049610.
 08-JUL-1997; 97US-0051926.
 16-JUL-1997; 97US-0052874.
 18-AUG-1997; 97US-0055724.
 22-AUG-1997; 97US-0056630.
 22-AUG-1997; 97US-0056631.
 22-AUG-1997; 97US-0056632.
 22-AUG-1997; 97US-0056636.
 22-AUG-1997; 97US-0056637.
 22-AUG-1997; 97US-0056662.
 22-AUG-1997; 97US-0056664.
 22-AUG-1997; 97US-0056845.
 22-AUG-1997; 97US-0056862.
 22-AUG-1997; 97US-0056864.
 22-AUG-1997; 97US-0056872.
 22-AUG-1997; 97US-0056874.
 22-AUG-1997; 97US-0056875.
 22-AUG-1997; 97US-0056876.
 22-AUG-1997; 97US-0056877.
 22-AUG-1997; 97US-0056878.
 22-AUG-1997; 97US-0056879.
 22-AUG-1997; 97US-0056880.
 22-AUG-1997; 97US-0056881.
 22-AUG-1997; 97US-0056882.
 22-AUG-1997; 97US-0056884.
 22-AUG-1997; 97US-0056886.
 22-AUG-1997; 97US-0056887.
 22-AUG-1997; 97US-0056888.
 22-AUG-1997; 97US-0056889.
 22-AUG-1997; 97US-0056892.
 22-AUG-1997; 97US-0056893.
 22-AUG-1997; 97US-0056894.
 22-AUG-1997; 97US-0056903.
 22-AUG-1997; 97US-0056908.
 22-AUG-1997; 97US-0056909.
 22-AUG-1997; 97US-0056910.
 22-AUG-1997; 97US-0056911.
 05-SEP-1997; 97US-0057650.
 05-SEP-1997; 97US-0057669.
 05-SEP-1997; 97US-0057761.
 12-SEP-1997; 97US-0058785.
 (HUMA-) HUMAN GENOME SCI INC.
 Bedharik DP, Brewer LA, Carter KC, Duan R, Ebner R, Endress GA,
 Feng P, Ferrie AM, Fischer CL, Florence KA, Greene JM, Hu JS,
 Kyaw H, Lafleur DW, Li Y, Moore PA, Ni J, Olsen HS, Rosen CA,
 Ruben SM, Shi Y, Soppet DR, Young PE, Yu GL, Zeng Z;
 WPI: 1998-506364/43.
 P-PSDB; W74815.
 New isolated human genes and the secreted polypeptide(s) they encode
 are useful for diagnosis and treatment of e.g. cancers, neurological
 disorders, immune diseases, inflammation or blood disorders
 Claim 1; Page 316-317; 721pp; English.

CC This sequence represents a nucleic acid molecule designated Gene 85 from
 CC the human cDNA clone HSDV29 (deposited as clone ATCC 209076) which
 CC encodes a secreted human protein. The gene can be used to generate
 CC fusion proteins by linking to the gene to a human immunoglobulin Fc
 CC portion (e.g. V59502) for increasing the stability of the fused protein
 CC as compared to the human protein only.
 CC The invention relates to 186 novel genes and their fragments (nucleic
 CC acid sequences: V59511-V59812; amino acid sequences W74731-W75026) which
 CC are useful for preventing, treating or ameliorating medical conditions
 CC e.g. by protein or gene therapy. Also, pathological conditions can be
 CC diagnosed by determining the amount of the new polypeptides in a sample
 CC or by determining the presence of mutations in the new polynucleotides.
 CC Specific uses are described for each of the 186 polynucleotides, based on
 CC which tissues they are most highly expressed in (see V59511 for described
 CC uses).
 XX
 SQ Sequence 2503 BP; 561 A; 705 C; 658 G; 568 T; 11 other;
 Query Match 0.6%; Score 17; DB 19; Length 2503;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2444 gctggcaggcgccctgg 2460
 |||||
 Db 583 gctggcaggcgccctgg 599
 RESULT 73
 Z42096
 ID Z42096 standard; cDNA; 2646 BP.
 XX
 AC Z42096;
 XX
 DT 31-JAN-2000 (first entry)
 XX
 DE Human endometrium tumour cDNA derived EST 116.
 XX
 KW Endometrium; human; tumour; cancer; anticancer; cytostatic; EST;
 KW treatment; uterine; gene therapy; expressed sequence tag; ss.
 XX
 OS Homo sapiens.
 XX
 PN DE19817948-A1.
 XX
 PD 21-OCT-1999.
 XX
 PF 17-APR-1998; 98DE-1017948.
 XX
 PR 17-APR-1998; 98DE-1017948.
 XX
 PA (META-) METAGEN GES GENOMFORSCHUNG MBH.
 XX
 PI Rosenthal A, Specht T, Hinzmann B, Schmitt A, Pilarsky C, Dahl E;
 XX
 DR WPI: 1999-591957/51.
 DR P-PSDB; Y60295, Y60296, Y60297.
 XX
 PT New nucleic acid sequences expressed in uterine cancer tissues, and
 PT derived polypeptides, for treatment of uterine and endometrial cancer
 PT and identification of therapeutic agents
 XX
 PS Claim 3; Page 253; 444pp; German.
 XX
 CC This invention describes novel human nucleic acid (cDNA) sequences (A),
 CC that are highly expressed in uterine tumour tissue and which have
 CC anticancer and cytostatic activity. (A) are used (i) for recombinant
 CC expression of polypeptides (B) and (ii) to isolate complete genes. (B)
 CC are used (i) to identify agents suitable for treatment of uterine or
 CC endometrial cancer; (ii) directly for treating these forms of cancer
 CC (including expression from gene therapy vectors) and (iii) for generation
 CC of specific antibodies. (A) are identified by assembling ESTs (expressed
 CC sequence tags) from a particular tissue type before comparison of

expression patterns. This allows a significantly longer fragment of the gene to be revealed, so should reduce the number of failures associated with the fact that ESTs from different libraries may represent different parts of the same unknown gene, distorting the estimated frequency of occurrence in a particular tissue. Z41981-Z42121 represent EST fragments derived from a human endometrium tumour cDNA library which encode the protein sequences represented in Y59941-Y60328.

Sequence 2646 BP; 787 A; 502 C; 546 G; 811 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 2646;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1463 agccccagcagagaaaa 1479
 |||||
 D 1717 agccccagcagagaaaa 1733

RESULT 74
 V68056
 V68056 standard; DNA; 2711 BP.
 V68056;
 02-FEB-1999 (first entry)
 Neurodegenerative polypeptide HHPDZ65 coding sequence.
 Neurodegenerative polypeptide: HHPDZ65; stroke; pain; epilepsy; therapy;
 neurodegenerative disease; ss.
 Homo sapiens.
 EP875570-A2.
 04-NOV-1998.
 15-APR-1998; 98EP-0302912.
 19-FEB-1998; 98GB-0003566.
 01-MAY-1997; 97GB-0008936.
 18-DEC-1997; 97EP-0310289.
 (SMIK) SMITHKLINE BEECHAM PLC.
 Bingham S, Davis J, Doe TR, Harrison DC, Topp S;
 WPI: 1998-559436/48.
 P-PSDB; W80315.
 HHPDZ65 polypeptide(s), their corresponding DNA, antibodies,
 agonists and antagonists - are useful in the treatment of stroke,
 pain, epilepsy and neurodegenerative diseases
 Claim 7; Page 14; 31pp; English.
 This sequence encodes the HHPDZ65 neurodegenerative polypeptide of the
 invention. HHPDZ65 is useful for the treatment of stroke, pain, epilepsy,
 neurodegenerative diseases and others. The DNAs and proteins are useful
 in a method for screening to identify compounds which stimulate or
 inhibit the function of the HHPDZ65 proteins. The polypeptides are useful
 in a process for diagnosing a disease or a susceptibility to a disease in
 a subject related to expression or activity of the HHPDZ65 polypeptides.
 Sequence 2711 BP; 559 A; 820 C; 852 G; 478 T; 2 other;

Query Match 0.6%; Score 17; DB 19; Length 2711;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2624 caccgtccccagagg 2640
 |||||
 Db 1879 caccgtccccagagg 1895

RESULT 75
 N81166/C
 ID N81166 standard; DNA; 2971 BP.
 XX
 AC N81166;
 XX
 DT 29-OCT-1990 (first entry)
 XX
 DE fdhF gene.
 XX
 KW repression/expression of foreign genes by oxygen/formate; ss.
 XX
 OS synthetic.
 XX
 FH Key Location/Qualifiers
 FT CDS 749..2896
 FT /*tag= a
 FT /product=fdhF
 FT RBS 737..740
 FT /*tag= b
 XX
 PN EP285152-A.
 XX
 PD 05-OCT-1988.
 XX
 PF 30-MAR-1988; 88EP-0105219.
 XX
 PR 31-MAR-1987; 87DE-3710633.
 PR 09-OCT-1987; 87DE-3735381.
 XX
 PA (BOEF) BOEHRINGER MANNHEIM GMBH.
 XX
 PI Birkmann A, Bock A;
 XX
 DR WPI: 1988-294149/42.
 XX
 PT New recombinant DNA contg consensus sequence and specific promoter -
 providing foreign gene repression under anaerobic conditions
 XX
 PS Disclosure; ; 12pp; German.
 XX
 CC Recombinant DNA contg a consensus sequence common to genes that are
 repressible by oxygen and inducible by formate under anaerobic
 CC conditions and the fdhF promoter used as a simple expression system
 CC for foreign genes. Expression does not require temp shifts or addition
 CC of inducers. (Formate is produced naturally by microorganisms in the
 CC late, anaerobic growth phase).
 CC See also N81165.
 XX
 SQ Sequence 2971 BP; 721 A; 771 C; 844 G; 635 T; 0 other;

Query Match 0.6%; Score 17; DB 9; Length 2971;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 112 ccatacgcagggcacc 128
 |||||
 Db 1781 CCATATCGCAGGCACC 1765

RESULT 76
 Q13115
 ID Q13115 standard; cDNA; 3000 BP.
 XX
 AC Q13115;
 XX
 DT 22-OCT-1991 (first entry)

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XX Encodes partial murine Natural Killer receptor.
XX NK; cytotoxic drugs; tumour cell; immunotherapy; mouse; ss.
XX Mus musculus.
XX Key Location/Qualifiers
XX sig_peptide 1..21
XX /tag= a
XX /note= "partial"
XX mat_peptide 22..2946
XX /tag= b
XX /product= murine NK receptor
XX
XX US7535206-A.
XX
XX 09-JUL-1991.
XX
XX 08-JUN-1990; 90US-0143578.
XX
XX 08-JUN-1990; 90US-0535206.
XX
XX (USSH ) NAT INST OF HEALTH.
XX
XX Ortaldo J, Young H, Anderson S;
XX
XX WPI; 1991-245694/33.
XX P-PSDB; R13320.
XX
XX DNA encoding a natural killer cell receptor - used to develop
XX prods. for the immuno-detection and immuno-therapy of tumours
XX
XX Disclosure; Fig 2; 30pp; English.
XX
XX Overlapping clones which make up this sequence were isolated from
XX a mouse Peripheral Blood Lymphocyte lambda gt10 cDNA library.
XX The protein encoded by this sequence is purified and can mediate
XX the cytolytic activity of mammalian cells. It specifically
XX distinguishes tumour cells making it a candidate for the development
XX of products for the immunodetection and immunotherapy of tumours.
XX See also Q13114.
XX
XX Sequence 3000 BP; 997 A; 661 C; 716 G; 626 T; 0 other;
XX
XX Query Match 0.6%; Score 17; DB 12; Length 3000;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 684 tttcagactccgagtc 700
XX |||||
XX 2743 tttcagactccgagtc 2759
XX
XX RESULT 77
XX V18471
XX V18471 standard; cDNA; 3156 BP.
XX
XX V18471;
XX
XX 14-SEP-1998 (first entry)
XX
XX T-cell surface antigen CD97 cDNA.
XX
XX T-cell surface antigen; CD97; human; inflammation; angiogenesis;
XX atherosclerosis; human; ss.
XX
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 49..2556
XX /tag= a

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FT sig_peptide 49..99
FT /tag= b
FT mat_peptide 100..2553
FT /tag= c
XX
XX WO9817796-A2.
XX
XX 30-APR-1998.
XX
XX 24-OCT-1997; 97WO-USI9772.
XX
XX 25-OCT-1996; 96US-0027871.
XX
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Kelly K;
XX
XX WPI; 1998-261492/23.
XX P-PSDB; W48756.
XX
XX New soluble CD97 alpha subunit isoform(s) - used to develop
XX products for the detection and treatment of inflammation,
XX atherosclerosis and angiogenesis
XX
XX Disclosure; Fig 1; 101pp; English.
XX
XX This polynucleotide comprises clone PAT276 that codes for human
XX T-cell surface antigen CD97 (see W48756). It was isolated from a
XX T-cell library enriched for mitogen-induced genes. The invention
XX relates to the previously unrecognised alpha subunit of CD97 that
XX acts in the establishment and maintenance of inflammation. Soluble
XX CD97 acts as an adhesion factor for endothelial cells and smooth
XX muscle cells, implicating it as a modulator of atherosclerosis.
XX CD97 alpha also acts as a motility factor to cells bearing the
XX alpha(V)beta3 receptor, indicative of a role in angiogenesis.
XX Soluble CD97 alpha1, alpha2, and alpha3 subunits (having different
XX combinations of EGF repeats) all originate as a proteolysin with the
XX beta subunit (see W48756). Host cells transfected with a nucleic
XX acid encoding a CD97 alpha subunit are claimed. CD97 alpha
XX subunit polypeptides, nucleic acids, antibodies and antagonists
XX (e.g. CD97 subunit antisense nucleic acids) are used in claimed
XX methods for: determining the degree of inflammation at a site;
XX identifying compounds that inhibit soluble CD97 alpha subunit
XX expression; inhibiting angiogenesis associated with chronic
XX inflammation; inhibiting atherosclerosis; and treating or inhibiting
XX CD97 associated inflammation.
XX
XX Sequence 3156 BP; 652 A; 973 C; 860 G; 671 T; 0 other;
XX
XX Query Match 0.6%; Score 17; DB 19; Length 3156;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1336 ccaggaggagtgccag 1352
XX |||||
XX 2683 ccaggaggagtgccag 2699
XX
XX RESULT 78
XX Z27969
XX ID Z27969 standard; DNA; 3156 BP.
XX
XX AC Z27969;
XX
XX 05-JAN-2000 (first entry)
XX
XX Human CD97 protein encoding DNA.
XX
XX Human; 7-transmembrane receptor; lectin-binding; mucin;
XX olfactomedin; cellular adhesion; atherosclerosis; gene therapy;
XX vascular disease; CD97; ss.
XX

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1S Homo sapiens.
2X W09945111-A1.
3X 10-SEP-1999.
4X 04-MAR-1999; 99WO-US04676.
5X 04-MAR-1998; 98US-0076782.
6X (ICOS-) ICOS CORP.
7X Hayflick JS;
8X WPI; 1999-571596/48.
9X P-PSDB; Y41090.
10X New human lectomedin receptor polypeptide, used to identify specific
11X binding partners for treating e.g. vascular disease
12X
13X Example 1; Page 106-110; 166pp; English.
14X
15X The invention provides purified and isolated human 7-transmembrane
16X receptor lectomedin polypeptide or its fragments. The lectomedin
17X polypeptide comprises extracellular lectin-binding, oifactomedin-like
18X and mucin-like domains. The polypeptide can be produced by standard
19X recombinant methodology. The polypeptide is involved in cellular adhesion
20X and cytoplasmic metabolic pathways that are modulated by extracellular
21X signaling. Specific binding to lectomedin-1 expressed on smooth muscle
22X cells may be required for proliferation of these cells in
23X atherosclerosis. The polypeptide is used to raise specific antibodies,
24X and to identify specific binding agents that modulate (increase or
25X decrease) its activity. The lectomedin nucleic acids are used as source
26X of probes and primers, and of therapeutic antisense, ribozyme or triplex-
27X forming agents, and in gene therapy to restore deficient lectomedin
28X activity. Specific binding agents of lectomedin are are used for treating
29X diseases that involve lectomedin activity, e.g. vascular diseases such as
30X atherosclerosis. The present sequence represents the DNA encoding the
31X human CD97 protein.
32X
33X Sequence 3156 BP; 651 A; 974 C; 860 G; 671 T; 0 other;
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XX 22-NOV-1994; 94US-0343760.
XX -F
XX 22-NOV-1994; 94US-0343760.
XX -R
XX (REGC ) UNIV CALIFORNIA.
XX -A
XX De Robertis EM, Sasai Y;
XX -I
XX WPI; 1997-525754/48.
XX -R
XX P-PSDB; W31559.
XX -P
XX DNA encoding Xenopus frog protein - that induces dorsal and neural
XX development and endodermal differentiation in vertebrates
XX -T
XX Claim 1; Columns 19-22; 27pp; English.
XX -S
XX This cDNA encodes a Xenopus protein "chordin". The functional recombinant
XX protein chordin has a defined sequence of 941 amino acids and can induce
XX dorsal and neural development and endodermal differentiation in
XX vertebrates. The presence of a hydrophobic signal sequence, four possible
XX N-glycosylation sites and conserved Cys-rich repeat regions suggest that
XX chordin is a secreted protein. The DNA sequence can be operatively linked
XX with an expression vector, to form a construct and a transformant can be
XX obtained by introducing the construct into a host. Chordin may be useful
XX as a component of culture media for culturing cells such as nerve or
XX muscle cells, for treating neurodegenerative diseases and damaged nerve
XX cells.
XX -C
XX Sequence 3796 BP; 1046 A; 841 C; 958 G; 951 T; 0 other;
XX -Q
Query Match 0.6%; Score 17; DB 18; Length 3796;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 270 ctctacgtcttctccga 286
QY 1036 ctctacgtcttctccga 1052
QY |||||
RESULT 81
N70558
D N70558 standard; DNA; 4216 BP.
XX AC
XX N70558;
XX -T
XX 29-APR-1991 (first entry)
XX -T
XX Sequence of nitrogen fixation gene H (nifH) promoter and coding
XX region and the nifH-nifD intergenic region.
XX -E
XX Rhizobium expression vector; plant expression vector;
XX -W Rhizobium; ss.
XX -X Rhizobium japonicum strain USDA 191.
XX -S
XX Key Location/Qualifiers
XX promoter 1921..1929
XX /tag= a
XX promoter 1933..1936
XX /tag= b
XX RBS 2012..2016
XX /tag= c
XX CDS 2024..2914
XX /tag= d
XX CDS 3011..4216
XX /tag= e
XX /label= nifD
XX misc_feature 1770..2023
XX /tag= f
XX /note= "claimed"
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XX EP211661-A.
XX PN
XX 25-FEB-1987.
XX PD
XX 07-AUG-1986; 86EP-0306105.
XX PF
XX 07-AUG-1985; 85US-0763800.
XX PR
XX (LUBR ) LUBRIZOL GENETICS I.
XX PA (LUBR-) LUBRIZOL GENETICS I.
XX PI
XX Appelbaum ER;
XX PI
XX WPI; 1987-051801/08.
XX DR
XX Nif promoter of fast-growing Rhizobium japonicum - used to drive
XX transcription in rhizobium of heterologous structural genes
XX PT
XX Disclosure; Fig 1; 37pp; English.
XX PS
XX Since the promoter region of the nifH operon has been isolated,
XX characterized and cloned, it is possible to delete the nifH and
XX nifHD nitrogenase genes and replace them with structural genes
XX isolated from an extraneous source. The extraneous genes thus placed
XX under the control of the nifH promoter can then be inserted into a
XX plasmid vector followed by conjugation into a fast-growing R.
XX japonicum strain.
XX CC
XX Sequence 4216 BP; 906 A; 1254 C; 1251 G; 793 T; 12 other;
XX -S
Query Match 0.6%; Score 17; DB 8; Length 4216;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1213 agctcaacctcaccac 1229
QY |||||
DB 3903 agctcaacctcaccac 3919
DB |||||
RESULT 82
T45351
ID T45351 standard; CDNA; 4258 BP.
XX AC
XX T45351;
XX -T
XX 18-MAR-1997 (first entry)
XX -T
XX Human colon carcinoma kinase 4 (CCK-4) CDNA.
XX DE
XX Colon carcinoma kinase 4; CCK-4; receptor tyrosine kinase;
XX KW signal transduction; colon cancer; diagnosis; gene therapy; ss.
XX -W
XX Homo sapiens.
XX -X
XX Key Location/Qualifiers
XX CDS 193..3405
XX /tag= a
XX sig_peptide 193..270
XX /tag= b
XX mat_peptide 271..3402
XX /tag= c
XX WO9637610-A2.
XX PN
XX 28-NOV-1996.
XX PD
XX 24-MAY-1996; 96WO-IB00696.
XX PF
XX 25-MAY-1995; 95US-0452630.
XX PR
XX (PLAC ) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.
XX PA
```

XX Alves F, Mossie K, Ullrich A;
XX WPI; 1997-021219/02.
XX P-PSDB; W08747.
XX New isolated receptor tyrosine kinase, CCK-4 - used for developing
XX prods. for the diagnosis and treatment of CCK-4 signal transduction
XX disorders, partic. colon cancer
XX Claim 1; Fig 1a-d; 129pp; English.
XX A cDNA sequence (T45351) codes for a novel human receptor tyrosine
XX kinase, colon carcinoma kinase-4 (W08747), or CCK-4, which is
XX preferentially expressed in cancerous colon tissue, compared with
XX normal colon. The cDNA sequence is a consensus of 6 overlapping
XX cDNA clones isolated from a human placenta library using primers
XX based on conserved motifs of protein tyrosine kinase catalytic
XX domains. CCK-4 nucleic acids can be used in the prodn. of
XX recombinant CCK-4 polypeptides, and as probes in the diagnosis and
XX screening of CCK-4 signal transduction disorders, esp. colon
XX cancer, and in gene therapy.
XX Sequence 4258 BP; 868 A; 1252 C; 1282 G; 856 T; 0 other;
XX
XX
XX Query Match 0.6%; Score 17; DB 18; Length 4258;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1338 agcaggagtgccag 1354
XX |||||
XX C 2648 agcaggagtgccag 2664
XX
XX RESULT 83
XX Q51806
XX ID 251806 standard; cDNA; 4308 BP.
XX
XX Q51806;
XX
XX 04-JUL-2000 (first entry)
XX Full length expanded cDNA sequence for human BAG-5 protein.
XX Human BAG-5; Bcl-2 associated athanogene-5; apoptosis; cell migration;
XX tumour cell metastasis; Hsc70/Hsp70-regulating protein; metastasis;
XX tumour cell proliferation; steroid hormone receptor function; ss.
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 247..1590
XX /*tag= a
XX /product= "Human BAG-5 protein"
XX
XX WO200014106-A1.
XX
XX 16-MAR-2000.
XX
XX 09-SEP-1999; 99WO-US21053.
XX
XX 09-SEP-1998; 98US-0150489.
XX
XX (BURN-) BURNHAM INST.
XX Reed JC, Takayama S;
XX WPI; 2000-256937/22.
XX P-PSDB; Y70517.
XX
XX BAG-1 related proteins from humans, Caenorhabditis elegans and
XX Schizosaccharomyces pombe useful for modulating tumor cell

PT proliferation, cell migration and metastasis and steroid hormone
PT receptor function -
XX Claim 12; Fig 17; 132pp; English.
XX
XX The present expanded cDNA sequence encodes human BAG-5 (Bcl-2 associated
XX athanogene-5) protein. BAG is a Hsc70/Hsp70-regulating protein
XX (Hsc70/Hsp70) is a molecular chaperone that participates in controlling
XX protein bioactivity, degradation, complex assembly/disassembly and
XX translocation across membranes). It competes with Hip for binding to the
XX Hsc70/Hsp70 ATPase binding domain and promotes substrate release. Gene
XX transfection studies indicate that BAG proteins influence a wide variety
XX of cellular phenotypes through their interactions with Hsc70/Hsp70,
XX including increasing resistance to apoptosis, promoting cell
XX proliferation, enhancing tumour cell migration and metastasis and
XX altering transcriptional activity of steroid hormones. BAG also
XX stimulates Hsc70-mediated adenosine triphosphate (ATP) hydrolysis by
XX accelerating ADP/ATP exchange.
XX Sequence 4308 BP; 1201 A; 897 C; 987 G; 1222 T; 1 other;
XX
XX
XX Query Match 0.6%; Score 17; DB 21; Length 4308;
XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 2126 ttgtgaaggaggagcag 2142
XX |||||
XX DB 1017 ttgtgaaggaggagcag 1033
XX
XX RESULT 84
XX Q51426
XX ID Q51426 standard; cDNA; 4488 BP.
XX
XX Q51426;
XX
XX 20-MAY-1994 (first entry)
XX Human FACC cDNA clone #1.
XX
XX Fanconi Anemia Group C; FACC; complementing cDNA; variant;
XX open reading frame; diagnosis; Fanconi anemia; ss.
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 174..1850
XX /*tag= a
XX /product= Human FACC
XX
XX polyA_signal 2240..2245
XX /*tag= b
XX polyA_signal 3042..3047
XX /*tag= c
XX repeat_unit 3163..3175
XX /*tag= d
XX /rpt_type= OTHER
XX /note= "palindrome"
XX repeat_unit 3289..3322
XX /*tag= e
XX /rpt_type= TANDEM
XX repeat_unit 3323..4455
XX /*tag= f
XX /rpt_type= TANDEM
XX
XX WO932435-A.
XX
XX 11-NOV-1993.
XX
XX 27-APR-1993; 93WO-CA00178.
XX
XX 29-APR-1992; 92US-0876285.
XX 21-JUL-1992; 92US-0918313.
XX
XX PR

15-JAN-1993; 93US-0003963.
 (HOSP-) HOSPITAL FOR SICK CHILDREN.
 (UNME-) UNITED MEDICAL & DENTAL SCHOOL GUYS.
 Buchwald M, Mathew CG, Strathdee CA, Wevrick R;
 WPI: 1993-368794/46.
 P-PSDB; R44139.
 Human cDNA which complements Fanconi Anaemia gp. C - used to develop prods. for use in diagnosis, study and therapy of Fanconi Anaemia
 Claim 1; Page 97-101; 137pp; English.
 The sequences given in Q51426-28 represent cDNA variants from the Fanconi Anemia Group C Complementing (FACC) cDNA. These three cDNA molecules are cellular variants of a single cDNA transcribed from the same gene. The three cDNAs each contain an identical open reading frame encoding the FACC protein. The FACC protein may be used for the diagnosis and study of Fanconi anemia.
 Sequence 4488 BP; 1052 A; 1092 C; 1168 G; 1176 T; 0 other;
 Query Match 0.6%; Score 17; DB 14; Length 4488;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 2105 tgaagccaccctggaag 2121
 |||||
 Db 4189 tgaagccaccctggaag 4205
 |||||
 RESULT 85
 ID V33945 standard; cDNA; 4567 BP.
 AC V33945;
 15-FEB-1999 (first entry)
 Fanconi anaemia complementation group C (FAC) cDNA.
 Fanconi anaemia complementation group C; FAC; apoptosis; haematopoiesis; bone marrow; chemotherapy; gene therapy; human; ds.
 Homo sapiens.
 Key Location/Qualifiers
 CDS 256..1929
 /*tag= a
 W09851792-A1.
 19-NOV-1998.
 15-MAY-1998; 98WO-US09975.
 15-MAY-1997; 97US-0046546.
 (BGHM) BRIGHAM & WOMENS HOSPITAL.
 Yousseoufian H;
 WPI: 1999-009774/01.
 P-PSDB; W68546.
 New conjugate of Fanconi anaemia molecule and peptide selective for haematopoietic precursor cells - inhibits apoptosis of these cells, for treating Fanconi anaemia and patients undergoing high-dose chemotherapy for cancer

XX Claim 6; Page 40-45; 72pp; English.
 PS This cDNA clone includes a coding region for human Fanconi anemia complementation group C (FAC, see W68546), a protein that modulates apoptosis in haematopoietic progenitor cells (HPC). The invention provides conjugates, including fusion proteins, comprising FAC and a targeting molecule which binds to a cell surface protein of the HPC and is internalised. Such targeting molecules include interleukin-3 (see W68547) and antibodies which recognise CD33 (see W68548-49). The conjugate, or a nucleic acid encoding it, can be used to deliver FAC to an HPC, specifically to inhibit apoptosis, particularly in patients exposed to high doses of chemotherapy for treatment of non-myeloid cancers, also to treat Fanconi anemia (by complementation of the genetic defect). Treatment of HPC is done in vitro, ex vivo (e.g. for recombinant production of conjugate in cell cultures) or in vivo. Treatment with FAC may eliminate the need for extensive bone marrow transplants to restore haematopoiesis after chemotherapy.
 Sequence 4567 BP; 1080 A; 1135 C; 1177 G; 1175 T; 0 other;
 Query Match 0.6%; Score 17; DB 20; Length 4567;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 2105 tgaagccaccctggaag 2121
 |||||
 Db 4268 tgaagccaccctggaag 4284
 |||||
 RESULT 86
 ID Q32352 standard; DNA; 5674 BP.
 AC Q32352;
 22-APR-1993 (first entry)
 MAGE-1 nucleic acid.
 melanoma antigen; MAGE TRA; melanoma antigen tumor rejection antigen; tumor rejection antigen precursor; MAGE; antigen E; gene family; ss.
 Homo sapiens.
 Key Location/Qualifiers
 CDS 3881..4711
 /*tag= a
 W09220356-A.
 26-NOV-1992.
 22-MAY-1992; 92WO-US04354.
 23-MAY-1991; 91US-0705702.
 09-JUL-1991; 91US-0728838.
 23-SEP-1991; 91US-0764364.
 12-DEC-1991; 91US-0807043.
 (LUDW-) LUDWIG INST CANCER RES.
 Boon T, Chomez P, De Plaen E, Lurquin C, Traversari C; Van Den Eynde B, Van Der Bruggen P, Van Pel A;
 WPI: 1992-415460/50.
 Nucleic acid mol. encoding a human tumour rejection antigen precursor - useful as an immunostimulant in a vaccine for treating and preventing cancers, also useful in diagnosis

S Disclosure; Page 71-73; 142pp; English.
 X The sequences given in Q32352-69 represent a new family of genes
 C referred to as melanoma antigens (MAGE). The cDNAs of this gene
 C family were identified during the isolation of the antigen E gene.
 C The MAGE cDNAs, when tested, did not transfer expression of antigen
 C E, but they did show substantial homology to the antigen E cDNA
 C sequence. The MAGE DNAs share a certain degree of homology with each
 C other and are expressed in tumour cells including several types of
 C human tumor cells as well as in human tumors. MAGE expression is not
 C restricted to melanomas. MAGE refers to a family of tumor rejection
 C antigen precursors. The antigens resulting from these genes are
 C referred to as MAGE TRAs or melanoma antigen tumor rejection antigens.
 C See also Q32351.

Sequence 5674 BP; 1277 A; 1644 C; 1568 G; 1185 T; 0 other;

Query Match 0.6%; Score 17; DB 13; Length 5674;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02; Indels 0; Gaps 0;
 Matches 17; Conservative 0; Mismatches 0;

Y 1652 ggtcctggggcaccctgg 1668
 |||||
 b 3994 ggtcctggggcaccctgg 4010

RESULT 87

Q72477 Q72477 standard; DNA; 5674 BP.

Q72477;

22-JUN-1995 (first entry)

Tumour rejection antigen MAGE-1 encoding DNA.

Tumour rejection antigen; melanoma antigen-1; MAGE-1; MAGE-3;
 cancer; cytolytic T cells; antigen D; human leucocyte antigen;
 ss.

Homo sapiens.

Key Location/Qualifiers
 CDS 3881..4711
 /*tag= a

W09423031-A.

13-OCT-1994.

17-MAR-1994; 94WO-US02877.

26-MAR-1993; 93US-0037230.

(LUDW-) LUDWIG INST CANCER RES.

Boon-falleur T, Gaugler B, Van Den EYNDE B, Van DER BRUGGEN P;

WPI; 1994-333192/41.

New tumour rejection antigen precursor MAGE3 - useful in
 treatment and diagnosis of cancer

Example 26; Page 59; 105pp; English.

Q72477 is the DNA sequence which encodes melanoma antigen-1
 (MAGE-1). Another melanoma antigen MAGE-3 is encoded by Q72470,
 this is a tumour rejection antigen precursor. Melanomas
 characterised by the expression of MAGE-3 can be detected, or
 monitored, by contacting a test sample with an agent that can
 recognise MAGE-3. The melanoma can be treated by the administration
 of cytolytic T cells specific for the complex of antigen D (the

CC mature rejection antigen derived from MAGE-3) and a human leucocyte
 CC antigen (esp. HLA-A1).

SQ Sequence 5674 BP; 1276 A; 1644 C; 1569 G; 1185 T; 0 other;

Query Match 0.6%; Score 17; DB 15; Length 5674;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02; Indels 0; Gaps 0;
 Matches 17; Conservative 0; Mismatches 0;

QY 1652 ggtcctggggcaccctgg 1668
 |||||
 Db 3994 ggtcctggggcaccctgg 4010

RESULT 88

X84113 X84113 standard; DNA; 5674 BP.

XX X84113;

DT 08-SEP-1999 (first entry)

DE MAGE-1 gene.

KW Tumour rejection antigen; vaccine; cancer; MAGE-1 gene; ss.

OS Homo sapiens.

PN US5925729-A.

PD 20-JUL-1999.

PF 02-MAY-1994; 94US-0142368.

PR 02-MAY-1994; 94US-0142368.

PR 23-MAY-1991; 91US-0705702.

PR 09-JUL-1991; 91US-0728838.

PR 23-SEP-1991; 91US-0764365.

PR 12-DEC-1991; 91US-0807043.

PA (LUDW-) LUDWIG INST CANCER RES.

PI Boon T, Chomez P, De Plaen E, Lurquin C, Traversari C;

PI Van Den Eynde B, Van Der Bruggen P, Van Pel A;

XX WPI; 1999-418294/35.

XX New tumour rejection antigen is useful as a vaccine against
 PT cancerous diseases

PS Disclosure; Column 39-46; 58pp; English.

CC This sequence represents the MAGE-1 gene sequence.

CC The invention relates to a tumour rejection antigen sequence that is
 CC useful as a tumour rejection antigen for vaccination against cancerous
 CC conditions.

SQ Sequence 5674 BP; 1276 A; 1644 C; 1569 G; 1185 T; 0 other;

Query Match 0.6%; Score 17; DB 20; Length 5674;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctggggcaccctgg 1668
 |||||
 Db 3994 ggtcctggggcaccctgg 4010

RESULT 89

T42117 T42117 standard; cDNA; 5720 BP.

conflict 5101 /tag= as /note= "C in genomic sequence"
conflict 5104..5114 /tag= at /note= "TGTAATTAGTG in genomic sequence"
conflict 5143 /tag= au /note= "A in genomic sequence"
conflict 5147 /tag= av /note= "A in genomic sequence"
conflict 5151..5169 /tag= aw /note= "GTGGCCCCCTCCCTCCCTCCTCAT in genomic sequence"
conflict 5247..5253 /tag= ax /note= "Deleted in genomic DNA"
conflict 5646..5663 /tag= ay /note= "AAAGCAAATTAATAAT in genomic sequence"
polyA_signal 5657..5663 /tag= az
WO9630402-A1.
03-OCT-1996.
26-MAR-1996; 96WO-US04101.
27-MAR-1995; 95US-041111.
(UYA) UNIV YALE.
Tao W, Wang W, Xu T, Yu W, Zhang S;
WPI: 1996-455275/45.
P-PSDB; W05177.
New isolated large tumour suppressor gene - used to develop prods.
for inhibiting cell proliferation or for enhancing proliferation
Disclosure; Page 109-115; 215pp; English.
This sequence encodes the Drosophila melanogaster large tumour
suppressor lats protein, and is a composite of 2 cDNAs (an initial
fragment from cDNA-9 and the rest from cDNA-A2). The sequence has
been isolated from a total imaginal disc cDNA library. The
corresponding genomic sequence is identical, except for 34 minor
differences, and has 7 introns. Two consensus polyadenylation sites
are present. A 141-bp sequence at the 3'-end of the lats transcript
is identical to the 5'-end of the untranslated sequence of the
Query Match 0.6%; Score 17; DB 17; Length 5720;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
814 tgggtctcaagcaag 830
|||||
4820 tgggtctcaagcaag 4836
RESULT 90
51508
251508 standard; DNA; 5720 BP.
251508;
21-JUN-2000 (first entry)
Drosophila melanogaster lats (large tumour suppressor) DNA.

KW Fruit fly; Lats; large tumour suppressor; cytostatic; vulnary;
KW cell overproliferation inhibitor; cdc2; cell cycle-dependent kinase;
KW treatment; screening; cancer; skin; ovarian tumour;
KW soft tissue sarcoma; pituitary disorder; gene therapy; hyperplasia;
KW LH; luteinizing hormone hypogonadotropic hypogonadism; metaplasia;
KW dysplasia; degenerative disorder; growth deficiency; physical trauma;
KW hypoproliferative disorder; lesion; wound; lats knock-out mouse; ds.
XX
OS Drosophila melanogaster.
XX
FH Key Location/Qualifiers
CDS 1103..4402
FT /tag= a
FT /product= "Lats protein"
FT polyA_signal 4555..4660
FT /tag= b
FT misc_feature 5013..5142
FT /tag= c
FT /note= "This region is identical to the 1-141
nucleotides of Drosophila plc-21 transcript"
XX
PN WO200010602-A1.
XX
PD 02-MAR-2000.
XX
XX 18-AUG-1999; 99WO-US19068.
XX 18-AUG-1998; 98US-0096996.
PR 18-AUG-1998; 98US-0096997.
XX
XX (UYA) UNIV YALE.
XX
XX Xu T, Tao W, St John MAR, Fei X, Fukumoto RK, Zhang S;
PI Turenchaik GS, Stewart RA;
XX
DR WPI: 2000-246496/21.
DR P-PSDB; Y70393.
XX
XX Use of lats proteins, complexes of lats and cdc2 for treating cancer
PT that is refractory to treatment by standard chemotherapy and radiation
PT therapy, and disorders associated with aberrant levels of cdc2 activity
PT
XX
PS Claim 44; Fig 15; 134pp; English.
XX
CC The present sequence is a DNA encoding Drosophila lats (large tumour
CC suppressor) protein which is a cell overproliferation inhibitor and a
CC negative regulator of cell cycle-dependent kinase cdc2/cyclin A.
CC The present sequence is useful for treating cancer that is refractory
CC to standard chemotherapy or radiation therapy such as hyperplasia,
CC metaplasia, or dysplasia, and disorders associated with aberrant
CC levels of cdc2 activity. Conditions treated by promoting cdc2 function
CC include degenerative disorders, growth deficiencies, hypoproliferative
CC disorders, physical trauma, lesions, and wounds. An animal model
CC preferably a mouse, in which a lats gene has been disrupted by homologous
CC recombination, e.g. a lats knock-out mouse, is used for screening
CC compounds that can be used to treat or prevent cancer, particularly
CC skin cancer, soft tissue sarcomas and ovarian tumours, and disorders
CC associated with pituitary dysfunction e.g. luteinizing hormone (LH)
CC hypogonadotropic hypogonadism. The lats DNA is also used in gene therapy.
XX
SQ Sequence 5720 BP; 1684 A; 1491 C; 1457 G; 1088 T; 0 other;
Query Match 0.6%; Score 17; DB 21; Length 5720;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
814 tgggtctcaagcaag 830
|||||
4820 tgggtctcaagcaag 4836
Query Match 0.6%; Score 17; DB 21; Length 5720;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
814 tgggtctcaagcaag 830
|||||
4820 tgggtctcaagcaag 4836

RESULT 91
 398902
 DT Q98902 standard; DNA; 5724 BP.
 XX
 AC Q98902;
 XX
 DT 28-FEB-1996 (first entry)
 XX
 DE Tumour rejection antigen (MAGE-1) gene.
 XX
 KW Tumour rejection antigen; MAGE-1; monoclonal antibody; MAb;
 XX diagnosis; immunoassay; cancer; ss.
 XX
 DS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 CDS 3881..4711
 FT /*tag= a
 FT /product= Tumour rejection antigen MAGE-1.
 FT /note= "The CDS is not indicated in the text of the
 FT specification but is suggested in the layout
 FT of the sequence."
 XX
 XX WO9520974-A1.
 XX
 XX 10-AUG-1995.
 XX
 XX 05-JAN-1995; 95WO-US000095.
 XX
 XX 01-FEB-1994; 94US-0190411.
 XX
 XX (LUDWIG) LUDWIG INST CANCER RES.
 XX (SLOK) SLOAN KETTERING INST CANCER RES.
 XX (SLOK) MEMORIAL SLOAN-KETTERING CANCER CENT.
 XX
 XX Boon-falleur T, Chen Y, Garin-chesa P, Old LJ; Rettig WJ;
 XX Stockert E, Van der bruggen P;
 XX WPT; 1995-283606/37.
 XX
 XX New monoclonal antibody binding specifically to MAGE-1 - useful for
 XX diagnosis and monitoring of cancer, also new hybridomas, recombinant
 XX MAGE-1 and immunogenic peptide(s)
 XX
 XX Disclosure; Page 16-19; 33pp; English.
 XX
 XX A monoclonal antibody directed against the tumour rejection antigen
 XX (MAGE-1) can be used to detect MAGE-1 in samples by standard
 XX immunoassay methods for diagnosis and monitoring of cancer etc. The
 XX monoclonal antibody is designated MA454 and is produced by the
 XX hybridoma deposited as ATCC HB11540. The monoclonal antibody is
 XX specific for MAGE-1, having no reactivity for MAGE-2 or MAGE-3.
 XX Peptide fragments of MAGE-1 (See R80518-20) may be useful as
 XX immunogens for production of the monoclonal antibody and antisera.
 XX
 XX Sequence 5724 BP; 1282 A; 1653 C; 1589 G; 1200 T; 0 other;

Query Match 0.6%; Score 17; DB 16; Length 5724;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 YY 1652 ggtctctgggacccctgg 1668
 ||||||||||||||||
 YY 3994 ggtctctgggacccctgg 4010

RESULT 92
 48772
 DT Q48772 standard; DNA; 6464 BP.
 XX
 AC Q48772;
 XX

DT 21-APR-1994 (first entry)
 XX
 DE Arylamine N-acetyl-transferase type 3.
 XX
 KW Arylamine N-acetyl-transferase; NAT; polymorphism; diagnosis;
 KW amino; aromatic substance; ds.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 CDS 723..1595
 FT /*tag= a
 FT exon 717..1936
 FT /*tag= b
 FT polyA_signal 1794..1799
 FT /*tag= c
 FT polyA_signal 1800..1805
 FT /*tag= d
 XX
 PN EP562547-A.
 XX
 XX 29-SEP-1993.
 XX
 XX 23-MAR-1993; 93EP-0104753.
 XX
 XX 23-MAR-1992; 92JP-0064669.
 XX
 XX (SARA) OTSUKA PHARM CO LTD.
 XX (TOKM-) TOKYO METROPOLITAN INST. NEUROSCIENCES.
 XX (TOKY-) ZH TOKYO SHINKEIKAGAKA SOGO KENKYUSHO.
 XX
 XX Deguchi T, Katsuragi K, Kinoshita M, Shin S;
 XX WPI; 1993-304947/39.
 XX P-PSDB; R41246.
 XX
 XX Polymorphic human arylamine N-acetyl-transferase genes - used to
 XX diagnose adverse effects caused by amino-contg. aromatic
 XX substances
 XX
 XX Claim 6; Page 37-41; 54pp; English.
 XX
 XX Sequences of three polymorphic human NAT genes are given in Q48767-
 XX Q48772. Detection of polymorphic human NAT genes allows diagnosis
 XX of adverse effects to be caused by amino-contg. aromatic substances.
 XX
 XX Sequence 6464 BP; 2087 A; 1189 C; 1282 G; 1906 T; 0 other;
 XX
 XX
 XX Query Match 0.6%; Score 17; DB 14; Length 6464;
 XX Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 YY 1032 aggtaccacgaagagc 1048
 ||||||||||||||||
 YY 5916 aggtaccacgaagagc 5932
 XX
 XX
 XX RESULT 93
 V52165
 ID V52165 standard; DNA; 10240 BP.
 XX
 AC V52165;
 XX
 XX 23-OCT-1998 (first entry)
 XX
 DE Streptococcus pneumoniae genome fragment SEQ ID NO:32.
 XX
 KW Streptococcus pneumoniae; S. pneumoniae; genome; diagnosis; assay;
 KW computer readable medium; vaccine; pharmaceutical composition; ds.
 XX
 XX Streptococcus pneumoniae.
 XX

```
XX WO9818931-A2.
XX
XX 07-MAY-1998.
XX
XX 30-OCT-1997; 97WO-US19588.
XX
XX 31-OCT-1996; 96US-0029960.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Barash SC, Choi GH, Dallon PJ, Dougherty BA, Fannon M;
XX Kunsch CA, Rosen CA;
XX WPI; 1998-272225/24.
XX
XX Computer-readable medium with recorded Streptococcus pneumoniae
XX polynucleotide sequences - useful in diagnostic kits and assays, and
XX pharmaceutical compositions and vaccines for Streptococcus
XX pneumoniae
XX
XX Claim 1; Page 326-332; 1409pp; English.
XX
XX The present invention describes a computer readable medium which has
XX the nucleotide sequences SEQ ID NO:1 to 391 (V52134 to V52524) recorded
XX on it, or a representative fragment of a sequence at least 95% identical
XX to SEQ ID NO:1 to 391. The nucleotide sequences depicted in SEQ ID NO:1
XX to 391 (V52134 to V52524) are genomic fragments from Streptococcus
XX pneumoniae. The present invention also describes an isolated nucleic acid
XX molecule encoding a homologue of any of the fragments of the S.pneumoniae
XX genome (SEQ ID NO:1 to 391) where the nucleic acid molecule is produced
XX by a process comprising: (a) screening a genomic DNA library using as a
XX probe a target sequence defined by any of the sequences in SEQ ID NO:1
XX to 391, identifying members of the library which contain sequences
XX that hybridise to the target sequence and isolating the nucleic acid
XX molecules from the members; or (b) isolating mRNA, DNA or cDNA produced
XX from an organism, amplifying nucleic acid molecules whose nucleotide
XX sequence is homologous to amplification primers derived from the
XX fragment of the S. pneumoniae genome to prime the amplification and
XX isolating the amplified sequences. The computer readable medium can be
XX used in a computer-based system for identifying fragments of the
XX S. pneumoniae genome of commercial importance, or expression modulating
XX fragments of the S. pneumoniae genome. Products from the present
XX invention can be used in diagnosis kits and assays, and pharmaceutical
XX compositions and vaccines for S. pneumoniae.
XX
XX Sequence 10240 BP; 2865 A; 1914 C; 2390 G; 3068 T; 3 other;

Query Match 0.6%; Score 17; DB 19; Length 10240;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1947 gatttgaagaggttca 1963
D 7643 gatttgaagaggttca 7659
|||||
RESULT 94
D20553
D X20553 standard; DNA; 10461 BP.
X20553;
X 05-MAY-1999 (first entry)
X
X Polynucleotide sequence from the genome of Treponema pallidum.
X Treponema pallidum infection; syphilis; Borrelia infection; animal;
X enzyme production; ds.
X Treponema pallidum.
X WO9859034-A2.
```

```
XX 30-DEC-1998.
XX
XX 23-JUN-1998; 98WO-US13041.
XX
XX 24-JUN-1997; 97US-0050667.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Fraser CM;
XX WPI; 1999-081273/07.
XX
XX New isolated Treponema pallidum nucleic acids - used to develop
XX products for the detection, diagnosis, characterisation, prevention
XX and therapy of T. pallidum infections, particularly syphilis
XX
XX Claim 1; Page 491-497; 1150pp; English.
XX
XX X20500-21243 represent polynucleotide sequences from the genome of
XX Treponema pallidum. The sequences can be used for detection,
XX diagnosis, characterisation, prevention and therapy for T. pallidum
XX infections, particularly syphilis. They can also be used for detecting
XX diseases related to Borrelia infections in animals, and for the
XX production of biosynthetic products such as enzymes.
XX
XX Sequence 10461 BP; 2325 A; 3411 C; 2493 G; 2213 T; 19 other;

Query Match 0.6%; Score 17; DB 20; Length 10461;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 161 ggggcacctgcgcacgc 177
DB 6768 ggggcacctgcgcacgc 6784
|||||
RESULT 95
V74675
ID V74675 standard; DNA; 10813 BP.
XX
XX V74675;
XX
XX 16-MAR-1999 (first entry)
XX
XX Staphylococcus aureus contig SEQ ID #364.
XX
XX Computer readable medium; vaccine; S.aureus infection; immunodetection;
XX cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;
XX skin infection; surgical wound infection; scalded skin syndrome;
XX toxic shock syndrome; ds.
XX
XX Staphylococcus aureus.
XX
XX Key Location/Qualifiers
XX misc_feature 301..360
XX /*tag= a
XX /*note= "these bases represent a line of missing text in
XX the sequence listing in the specification. They
XX are included to maintain the nucleotide numbering
XX given in the specification for this DNA sequence"
XX
XX misc_feature 2101..2160
XX /*tag= b
XX /*note= "these bases represent a line of missing text in
XX the sequence listing in the specification. They
XX are included to maintain the nucleotide numbering
XX given in the specification for this DNA sequence"
XX
XX misc_feature 3901..3960
XX /*tag= c
XX /*note= "these bases represent a line of missing text in
XX the sequence listing in the specification. They
XX are included to maintain the nucleotide numbering
XX given in the specification for this DNA sequence"
XX
```

```

I  misc_feature 5701..5760 given in the specification for this DNA sequence"
II /tag= d
III /note= "these bases represent a line of missing text in
IV the sequence listing in the specification. They
V are included to maintain the nucleotide numbering
VI given in the specification for this DNA sequence"
VII
VIII misc_feature 7501..7560
IX /tag= e
X /note= "these bases represent a line of missing text in
XI the sequence listing in the specification. They
XII are included to maintain the nucleotide numbering
XIII given in the specification for this DNA sequence"
XIV
XV misc_feature 9301..9360
XVI /tag= f
XVII /note= "these bases represent a line of missing text in
XVIII the sequence listing in the specification. They
XIX are included to maintain the nucleotide numbering
XX given in the specification for this DNA sequence"
XXI
XXII EP786519-A2.
XXIII
XXIV 30-JUL-1997.
XXV
XXVI 07-JAN-1997; 97EP-0100117.
XXVII
XXVIII 05-JAN-1996; 96US-0009861.
XXIX
XXX (HUMA-) HUMAN GENOME SCI INC.
XXXI
XXXII Barash SC, Choi GH, Dillon PJ, Fannon MR, Kunsch CA;
XXXIII Rosen CA;
XXXIV WPI; 1997-374922/35.
XXXV
XXXVI Polynucleotide(s) and proteins derived from Staphylococcus aureus -
XXXVII stored on computer readable medium and used in the production of
XXXVIII anti-S.aureus vaccines
XXXIX
XXXX Claim 1; Page 1248-1254; 3271pp; English.
XXXXI
XXXXII This sequence represents one of 5191 Staphylococcus aureus DNA sequences
XXXXIII of the invention. The DNA sequences are recorded on a computer readable
XXXXIV medium, preferably selected from a floppy or hard disk, random access
XXXXV memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
XXXXVI the S.aureus DNA sequences allows putative functions to be assigned so
XXXXVII that protein-encoding or regulatory regions of commercial, therapeutic or
XXXXVIII industrial importance can be obtained. Specifically, sequences which are
XXXXIX likely to encode antigens have been identified and these polypeptides can
XXXXX be used in a vaccine composition against S.aureus infection. The
XXXXXI polypeptides can also be used in a kit for the immunodetection of
XXXXXII S.aureus in a sample. S.aureus is implicated in numerous human diseases,
XXXXXIII including cellulitis, eyelid infections, food poisoning, osteomyelitis,
XXXXXIV skin and surgical wound infections, scalded skin syndrome, toxic shock
XXXXXV syndrome, etc. Organisms transformed with the DNA sequences can be used
XXXXXVI for recombinant production of the polypeptides. The new DNA sequences
XXXXXVII (and their fragments) are useful as primers or probes for isolating
XXXXXVIII homologues of any of the S.aureus DNA sequences contained on the
XXXXXIX computer readable medium.
XXXXXX
XXXXXXI Sequence 10813.BP; 3513 A; 1895 C; 1594 G; 3451 T; 360 other;
XXXXXXII
XXXXXXIII Query Match 0.6%; Score 17; DB 18; Length 10813;
XXXXXXIV Best Local Similarity 100.0%; Pred. No. 1.7e+02;
XXXXXXV Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XXXXXXVI
XXXXXXVII 474 ccacaactggaataa 490
XXXXXXVIII
XXXXXXIX db 10019 ccacaactggaataa 10035
XXXXXXX
XXXXXXXI RESULT 96

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Q70447
ID Q70447 standard; DNA; 11236 BP.
XX
XX AC Q70447;
XX
XX DT 21-MAR-1995 (first entry)
XX
XX DE Rat nestin gene - its product is useful to identify brain tumours.
XX
XX KW nestin gene; brain tumour; neoplastic cells; glial; neuronal;
XX muscle; neural multipotential stem cell; mammalian brain; detection;
XX diagnosis; medulloblastoma; glioblastoma; oligodendroglioma; ds.
XX
XX OS Rattus rattus.
XX
XX FH Key Location/Qualifiers
XX misc_feature 2463
XX FT /tag= a
XX FT /note= "start of primary transcript"
XX CDS 2589..10821
XX FT /tag= b
XX FT /product= Nestin protein
XX FT 3089..3090
XX FT /tag= c
XX FT /transl_except= pos:3087..3091, aa:His, Arg
XX FT /note= "sequence should be CAY CGG, ie. Y or C has
XX been deleted in the sequence given"
XX
XX FT misc_feature 3104
XX FT /tag= d
XX FT /note= "apparent inclusion of a nucleotide"
XX intron 3375..4339
XX FT /tag= e
XX FT 4340..4464
XX FT /tag= f
XX FT 4465..6119
XX FT /tag= g
XX FT 6120..6193
XX FT /tag= h
XX FT 6194..6388
XX FT /tag= i
XX FT 6389..10821
XX FT /tag= j
XX
XX US5338839-A.
XX
XX PD 16-AUG-1994.
XX
XX PF 12-APR-1988; 88US-0180548.
XX
XX PR 12-APR-1988; 88US-0180548.
XX PR 02-JUN-1988; 88US-0201762.
XX PR 25-OCT-1990; 90US-0603803.
XX PR 22-FEB-1991; 91US-0660412.
XX PR 19-MAR-1992; 92US-0853913.
XX
XX PA (MASI ) MASSACHUSETTS INST TECHNOLOGY.
XX
XX PI Lendahl U, McKay RDG;
XX
XX DR WPI; 1994-263332/32.
XX
XX DR P-PSDB; R60126.
XX
XX PT Nucleotide and protein sequences for human and rat nestin -
XX distinguishes neural multipotential stem cells and brain tumour
XX cells from more differentiated cell types; for use in the
XX diagnosis of brain tumours
XX
XX PS Claim 1; Column 25-34; 45pp; English.
XX
XX CC Q70447 is the rat nestin gene encoding nestin protein (R60126).
XX Nestin protein expression distinguishes neural multipotential stem
XX cells and brain tumour cells from the more differentiated neural
XX cell types (eg., neuronal, glial and muscle cells of the adult brain).
XX

```

CC The nestin protein can be used in diagnosing tumours of the brain,
 CC such as medulloblastomas, glioblastomas and oligodendroglioma.
 CC (See also Q70448).

XX Sequence 11236 BP; 2876 A; 2678 C; 3258 G; 2424 T; 0 other;

Query Match 0.6%; Score 17; DB 15; Length 11236;
 Best Local Similarity 100.0%; Pred. No. 1.7e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

NY 1877 ccttcaggaaagggctg 1893
 |||||
 3 3466 ccttcaggaaagggctg 3482

RESULT 97
 220535/c
 X20535 standard; DNA; 21170 BP.

XX X20535;

05-MAY-1999 (first entry)

Polynucleotide sequence from the genome of Treponema pallidum.

Treponema pallidum infection; syphilis; Borrelia infection; animal;
 enzyme production; ds.

Treponema pallidum.

WO9859034-A2.

30-DEC-1998.

23-JUN-1998; 98WO-US13041.

24-JUN-1997; 97US-0050667.

(HUMA-) HUMAN GENOME SCI INC.

Fraser CM;

WPI; 1999-081273/07.

New isolated Treponema pallidum nucleic acids - used to develop
 products for the detection, diagnosis, characterisation, prevention
 and therapy of T. pallidum infections, particularly syphilis

Claim 1; Page 389-401; 1150pp; English.

X20500-21243 represent polynucleotide sequences from the genome of
 Treponema pallidum. The sequences can be used for detection,
 diagnosis, characterisation, prevention and therapy for T. pallidum
 infections, particularly syphilis. They can also be used for detecting
 diseases related to Borrelia infections in animals, and for the
 production of biosynthetic products such as enzymes.

Sequence 21170 BP; 4629 A; 5015 C; 6107 G; 5390 T; 29 other;

Query Match 0.6%; Score 17; DB 20; Length 21170;
 Best Local Similarity 100.0%; Pred. No. 1.6e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

85 ccgacgcaccatgctg 101
 |||||
 3 1835 CCGACGCACCATGCTG 1819

RESULT 98
 23005
 X83005 standard; DNA; 29604 BP.

XX X83005;
 AC
 XX 31-AUG-1999 (first entry)
 XX
 DE Partial mouse WRN genomic sequence #1.
 XX
 KW Mouse; WRN; Werner's syndrome; detection; diagnosis; autosomal;
 KW recessive disorder; phenotype; ss.
 OS Mus musculus.

XX WO9724435-A1.

XX 10-JUL-1997.

XX 30-DEC-1996; 96WO-US20785.

XX 12-APR-1996; 96US-0632175.

XX 29-DEC-1995; 95US-0009409.

XX 29-DEC-1995; 95US-0580539.

XX 30-JAN-1996; 96US-0010835.

XX 30-JAN-1996; 96US-0594242.

(DARW-) DARWIN MOLECULAR CORP.
 (OSHI/) OSHIMA J.

Fu Y, Mulligan J, Oshima J, Schellenberg GD, Yu C;

WPI; 1997-363671/33.

Isolated nucleic acid molecule encoding the WRN gene product -
 useful for detection and treatment of Werner's syndrome, and related
 diseases

Claim 1; Fig 7; 153pp; English.

This sequence represents a fragment of the genomic sequence containing
 the coding region for the mouse WRN gene (X83004). The corresponding
 human gene (X83001) encodes a protein related to Werner's syndrome.
 The products can be used for the detection and treatment of Werner's
 syndrome (WS), an autosomal recessive disorder with a complex phenotype,
 as well as related diseases.

Sequence 29604 BP; 7634 A; 5861 C; 5985 G; 10123 T; 1 other;

Query Match 0.6%; Score 17; DB 18; Length 29604;
 Best Local Similarity 100.0%; Pred. No. 1.6e+02;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 525 aaaggaataagaactggc 541

Db 14826 aaaggaataagaactggc 14842

RESULT 99
 230163/c
 ID 230163 standard; DNA; 34094 BP.

XX X230163;

XX 26-JAN-2000 (first entry)

Complete nucleotide sequence of the PAV-3 genome.

XX PAV-3; defective recombinant PAV vector; live recombinant virus;
 KW subunit vaccine; nucleic acid immunisation; gene therapy;
 KW genetic disease; hemophilia; cystic fibrosis; cancer; viral infection;
 KW acquired immune deficiency syndrome; PAV antigen; porcine pathogen; ds.
 XX Porcine adenovirus Type 3.
 OS XX

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PN WO9953047-A2.
XX 21-OCT-1999.
XX 15-APR-1999; 99WO-US08220.
XX 15-APR-1998; 98US-0081882.
XX (UYSA-) UNIV SASKATCHEWAN.
XX Reddy PS, Tikoo SK, Babiuk LA;
XX WPI; 1999-620422/53.
XX New nucleic acids from the genome of porcine adenovirus-3, and derived
XX gene therapy vectors, particularly for immunization
XX Example 2; Fig 1; 87pp; English.
XX The present sequence represents the complete nucleotide sequence of the
XX genome of porcine adenovirus-3 (PAV-3). The specification also describes
XX a defective recombinant PAV vector comprising inverted terminal repeats
XX (ITR), packaging sequences and at least one heterologous nucleotide
XX sequence (II), but lacking E1 functions. The defective vectors replicate
XX inefficiently in cells (other than helper cells) so are unlikely to be
XX immunogenic. Deletion of the E1 (and optionally other regions) increases
XX the size of heterologous insert that can be packaged. The PAV-3
XX polynucleotides sequences are used to produce (recombinant or defective)
XX vectors that can express heterologous proteins, e.g. for making live,
XX recombinant virus or subunit vaccines, for nucleic acid immunisation or
XX for gene therapy (e.g. of genetic diseases such as hemophilia or cystic
XX fibrosis, cancer, or viral infections, including acquired immune
XX deficiency syndrome), also for in vitro expression of recombinant
XX antigens (for antibody production), antisense RNA, ribozymes or
XX therapeutic proteins. They are also used diagnostically to detect PAV
XX antigens and/or nucleic acid. The vectors may be used in human or
XX veterinary medicine, but particularly for expressing protective
XX determinants of porcine pathogens. Regulatory regions may be used to
XX control expression of heterologous genes. Antibodies raised against PAV-3
XX polypeptides can also be used for diagnosis (to detect PAV-specific
XX antigen).
XX Sequence 34094 BP; 6240 A; 11070 C; 10693 G; 6091 T; 0 other;
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XX Query Match 0.6%; Score 17; DB 20; Length 34094;
XX Best Local Similarity 100.0%; Pred. No. 1.6e+02;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 330 ctcacgagcagagcaca 346
XX |||||
XX 4686 CTCATGACGAGGACACAA 4670
XX
XX RESULT 100
XX 32020/C
XX 32020 standard; DNA; 38734 BP.
XX
XX 32020;
XX
XX 10-JAN-2000 (first entry)
XX
XX Human METH1 related EST AL021529.
XX
XX Human; METH1; METH2; anti-angiogenic; metalloprotease thrombospondin;
XX cancer; diagnosis; hyperproliferative disorder; autoimmune disease;
XX angiogenesis inhibitor; abnormal wound healing; inflammation;
XX rheumatoid arthritis; psoriasis; endometrial bleeding disorder;
XX diabetic retinopathy; macula degeneration; haemangioma; detection;
XX arterial-venous malformation; immune deficiency; SS.
XX
XX Homo sapiens.
```

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PN WO9937660-A1.
XX 29-JUL-1999.
XX 22-JAN-1999; 99WO-US01313.
XX 23-JAN-1998; 98US-0072298.
XX 28-AUG-1998; 98US-0098539.
XX (IRUE/) IRUELA-ARISPE L.
XX (HAST/) HASTINGS G A.
XX (RUBE/) RUBEN S M.
XX
XX Iruela-Arispe L, Hastings GA, Ruben SM;
XX WPI; 1999-590684/50.
XX New isolated metalloprotease thrombospondin polypeptides, useful for
XX treating hyperproliferative disorders, cancers or autoimmune disorders
XX
XX Disclosure; Page 296-321; 457pp; English.
XX
XX 232000 and 232001 encode, and Y49501 and Y49502 represent, human
XX metalloprotease thrombospondin (METH) proteins METH1 and METH2
XX respectively. METH1 and METH2 have been found to be potent inhibitors of
XX angiogenesis both in vitro and in vivo. They can be used for treating
XX cancer and other disorders related to angiogenesis including abnormal
XX wound healing, inflammation, rheumatoid arthritis, psoriasis,
XX endometrial bleeding disorders, diabetic retinopathy, some forms of
XX macula degeneration, haemangiomas, and arterial-venous malformations.
XX They may be useful in treating deficiencies or disorders of the immune
XX system, by activating or inhibiting the proliferation, differentiation,
XX or mobilisation (chemotaxis) of immune cells. The etiology of these
XX immune deficiencies or disorders may be genetic, somatic, such as
XX cancer or some autoimmune disorders, acquired (e.g. by chemotherapy or
XX toxins), or infectious. They can also be used to treat inflammatory
XX conditions, both chronic and acute conditions. The products can also be
XX used for detection and diagnosis. 232002 to 232080, and Y49503 to Y49511
XX represent sequences given in the exemplification of the present
XX invention.
XX
XX Sequence 38734 BP; 6142 A; 13140 C; 13585 G; 5867 T; 0 other;
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Query Match 0.6%; Score 17; DB 20; Length 38734;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 11 ggtgaccgagcgcttcc 27
|||
DB 37241 GGTGACCGCGGCTTTC 37225
Search completed: February 18, 2001, 16:04:40
Job time: 25989 sec
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GenCore version 4.5
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DM nucleic - nucleic search, using sw model

Run on: February 18, 2001, 07:28:55 ; Search time 96.47 Seconds
(without alignments)
4941.559 Million cell updates/sec

Title: US-09-434-382-3
Perfect score: 2958
Sequence: 1 cgcggcgtagtgaccggc.....aataaagattgattgcaa 2958

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0
Searched: 280836 seqs, 80580151 residues

Word size : 0

Total number of hits satisfying chosen parameters: 561672

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 100 summaries

Database : Issued Patents_NA.*
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2: /cgn1_7/ptodata/1/ina/5B_COMB.seq.*
3: /cgn1_7/ptodata/1/ina/6_COMB.seq.*
4: /cgn1_7/ptodata/1/ina/PCTUS_COMB.seq.*
5: /cgn1_7/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Length	ID	Description
1	19	0.6	2955	2	US-08-867-941-9
2	19	0.6	3000	2	US-08-867-941-8
3	19	0.6	7641	2	US-08-867-941-6
4	18	0.6	1152	3	US-09-045-186-1
5	18	0.6	1152	3	US-09-045-186-3
6	18	0.6	2186	2	US-08-878-546-9
7	18	0.6	2605	2	US-08-680-395-4
8	17	0.6	289	2	US-08-967-101-23
9	17	0.6	289	2	US-08-592-541-23
10	17	0.6	289	3	US-09-124-698-23
11	17	0.6	304	2	US-08-611-757-20
12	17	0.6	304	4	PCT-US95-05380-20
13	17	0.6	1084	2	US-08-184-009-110
14	17	0.6	1084	2	US-08-458-356-110
15	17	0.6	1094	2	US-08-184-009-109
16	17	0.6	1094	2	US-08-458-356-109
17	17	0.6	1691	2	US-08-993-118-8
18	17	0.6	1691	3	US-08-845-528C-8
19	17	0.6	1816	2	US-08-951-148-2
20	17	0.6	1816	2	US-09-165-234-2
21	17	0.6	1816	3	US-09-274-570-2
22	17	0.6	2419	1	US-07-807-043B-7
23	17	0.6	2419	1	US-08-299-849B-7
24	17	0.6	2419	2	US-08-142-368A-7
25	17	0.6	2419	3	US-08-967-727-7
26	17	0.6	2420	1	US-08-465-167A-23
27	17	0.6	3457	1	US-08-295-882-1
28	17	0.6	3796	1	US-08-343-760A-1

1	4488	0.6	17	29	US-08-441-430-1	Sequence 1, Appl
2	5674	0.6	17	30	US-07-807-043B-8	Sequence 8, Appl
3	5674	0.6	17	31	US-08-190-411A-1	Sequence 1, Appl
4	5674	0.6	17	32	US-08-299-849B-8	Sequence 1, Appl
5	5674	0.6	17	33	US-08-560-024-1	Sequence 1, Appl
6	5674	0.6	17	34	US-08-142-368A-8	Sequence 8, Appl
7	5674	0.6	17	35	US-08-967-727-8	Sequence 6, Appl
8	5674	0.6	17	36	US-08-321-478-6	Sequence 1, Appl
9	11236	0.6	17	37	US-07-853-913-1	Sequence 1, Appl
10	29604	0.6	17	38	US-08-781-891-207	Sequence 207, App
11	68750	0.6	17	39	US-09-335-409-1	Sequence 1, Appl
12	76	0.5	16	40	US-07-753-110B-12	Sequence 12, Appl
13	76	0.5	16	41	US-08-503-730-16	Sequence 16, Appl
14	76	0.5	16	42	US-08-507-634-13	Sequence 13, Appl
15	105	0.5	16	43	US-08-717-294-93	Sequence 93, Appl
16	233	0.5	16	44	US-08-687-080-70	Sequence 70, Appl
17	252	0.5	16	45	US-08-630-822A-97	Sequence 97, Appl
18	252	0.5	16	46	US-09-005-069-97	Sequence 97, Appl
19	252	0.5	16	47	US-08-906-769-104	Sequence 104, App
20	252	0.5	16	48	US-08-906-616-104	Sequence 104, App
21	252	0.5	16	49	US-08-817-795-104	Sequence 104, App
22	252	0.5	16	50	US-08-639-075A-104	Sequence 104, App
23	252	0.5	16	51	PCT-US95-14442A-104	Sequence 104, App
24	294	0.5	16	52	US-08-611-757-98	Sequence 98, Appl
25	294	0.5	16	53	PCT-US95-05980-98	Sequence 98, Appl
26	486	0.5	16	54	US-08-937-931-9	Sequence 9, Appl
27	489	0.5	16	55	US-08-334-254-7	Sequence 7, Appl
28	489	0.5	16	56	US-08-848-131-7	Sequence 7, Appl
29	575	0.5	16	57	PCT-US95-14792-7	Sequence 7, Appl
30	575	0.5	16	58	US-08-507-016-8	Sequence 8, Appl
31	578	0.5	16	59	PCT-US91-06418-4	Sequence 4, Appl
32	609	0.5	16	60	US-08-338-579A-94	Sequence 94, Appl
33	654	0.5	16	61	US-08-911-319A-2	Sequence 2, Appl
34	654	0.5	16	62	US-09-352-819-2	Sequence 2, Appl
35	714	0.5	16	63	US-07-789-738-3	Sequence 3, Appl
36	773	0.5	16	64	US-07-789-738-5	Sequence 5, Appl
37	808	0.5	16	65	US-08-651-136C-15	Sequence 15, Appl
38	815	0.5	16	66	US-08-906-769-128	Sequence 128, App
39	815	0.5	16	67	US-08-906-616-128	Sequence 128, App
40	815	0.5	16	68	US-08-639-075A-128	Sequence 128, App
41	1031	0.5	16	69	US-08-651-136C-19	Sequence 19, Appl
42	1048	0.5	16	70	US-08-651-136C-17	Sequence 17, Appl
43	1152	0.5	16	71	US-08-933-750C-81	Sequence 81, Appl
44	1152	0.5	16	72	US-09-234-613-81	Sequence 81, Appl
45	1225	0.5	16	73	US-08-739-485-4	Sequence 4, Appl
46	1329	0.5	16	74	US-08-360-758-1	Sequence 1, Appl
47	1333	0.5	16	75	US-08-889-425-3	Sequence 3, Appl
48	1341	0.5	16	76	US-09-032-372-9	Sequence 9, Appl
49	1362	0.5	16	77	US-08-374-155A-7	Sequence 7, Appl
50	1362	0.5	16	78	US-08-785-396-7	Sequence 7, Appl
51	1389	0.5	16	79	US-08-458-023B-1	Sequence 1, Appl
52	1389	0.5	16	80	US-09-111-556A-1	Sequence 1, Appl
53	1524	0.5	16	81	US-08-135-510-4	Sequence 4, Appl
54	1524	0.5	16	82	US-08-483-852-4	Sequence 4, Appl
55	1524	0.5	16	83	US-08-477-953-4	Sequence 4, Appl
56	1524	0.5	16	84	US-08-409-122-1	Sequence 1, Appl
57	1524	0.5	16	85	US-08-408-669-1	Sequence 1, Appl
58	1524	0.5	16	86	US-08-477-952-4	Sequence 4, Appl
59	1713	0.5	16	87	US-08-386-727-5	Sequence 5, Appl
60	1713	0.5	16	88	US-08-600-452A-5	Sequence 5, Appl
61	1713	0.5	16	89	US-08-467-948A-1	Sequence 1, Appl
62	1713	0.5	16	90	US-08-467-947A-1	Sequence 1, Appl
63	1721	0.5	16	91	US-07-688-352C-13	Sequence 13, Appl
64	1721	0.5	16	92	US-08-474-379C-13	Sequence 13, Appl
65	1721	0.5	16	93	US-09-146-249A-13	Sequence 13, Appl
66	1721	0.5	16	94	US-08-206-188B-13	Sequence 13, Appl
67	1721	0.5	16	95	PCT-US91-02714-13	Sequence 13, Appl
68	1766	0.5	16	96	US-08-481-814A-2	Sequence 2, Appl
69	1790	0.5	16	97	US-08-993-228-1	Sequence 1, Appl
70	1977	0.5	16	98	US-09-231-529-2	Sequence 2, Appl
71	1990	0.5	16	99	US-09-255-911-1	Sequence 1, Appl
72	2069	0.5	16	100	US-08-619-554-7	Sequence 7, Appl

ALIGNMENTS

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RESULT 1
US-08-867-941-9
Sequence 9, Application US/08867941
Patent No. 5977337
GENERAL INFORMATION:
APPLICANT: Loosmore, Sheena M
APPLICANT: Du, Run-Pan
APPLICANT: Wang, Quijun
APPLICANT: Yang, Yan-Ping
APPLICANT: Klein, Michel H
TITLE OF INVENTION: LACTOFERRIN RECEPTOR GENES OF MORAXELLA
NUMBER OF SEQUENCES: 67
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sim & McBurney
STREET: 6th Floor, 330 University Avenue
CITY: Toronto
STATE: Ontario
COUNTRY: Canada
ZIP: M5G 1R7
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/867,941
FILING DATE: 03-JUN-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Stewart, Michael I
REGISTRATION NUMBER: 24,973
REFERENCE/DOCKET NUMBER: 1038-681 MIS:jb
TELECOMMUNICATION INFORMATION:
TELEPHONE: (416) 595-1155
TELEFAX: (416) 595-1163
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 2955 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-867-941-9

Query Match 0.6%; Score 19; DB 2; Length 2955;
Best Local Similarity 100.0%; Pred. No. 7.5;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 811 tcttggtgctcaagcaaa 829
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D 591 TCTTGGTGTCTCAAGCAAA 609

RESULT 2
US-08-867-941-8
Sequence 8, Application US/08867941
Patent No. 5977337
GENERAL INFORMATION:
APPLICANT: Loosmore, Sheena M
APPLICANT: Du, Run-Pan
APPLICANT: Wang, Quijun
APPLICANT: Yang, Yan-Ping
APPLICANT: Klein, Michel H
TITLE OF INVENTION: LACTOFERRIN RECEPTOR GENES OF MORAXELLA
NUMBER OF SEQUENCES: 67
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sim & McBurney
STREET: 6th Floor, 330 University Avenue
CITY: Toronto
```

```
STATE: Ontario
COUNTRY: Canada
ZIP: M5G 1R7
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/867,941
FILING DATE: 03-JUN-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Stewart, Michael I
REGISTRATION NUMBER: 24,973
REFERENCE/DOCKET NUMBER: 1038-681 MIS:jb
TELECOMMUNICATION INFORMATION:
TELEPHONE: (416) 595-1155
TELEFAX: (416) 595-1163
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 3000 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-867-941-8

Query Match 0.6%; Score 19; DB 2; Length 3000;
Best Local Similarity 100.0%; Pred. No. 7.5;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 811 tcttggtgctcaagcaaa 829
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DB 636 TCTTGGTGTCTCAAGCAAA 654

RESULT 3
US-08-867-941-6
Sequence 6, Application US/08867941
Patent No. 5977337
GENERAL INFORMATION:
APPLICANT: Loosmore, Sheena M
APPLICANT: Du, Run-Pan
APPLICANT: Wang, Quijun
APPLICANT: Yang, Yan-Ping
APPLICANT: Klein, Michel H
TITLE OF INVENTION: LACTOFERRIN RECEPTOR GENES OF MORAXELLA
NUMBER OF SEQUENCES: 67
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sim & McBurney
STREET: 6th Floor, 330 University Avenue
CITY: Toronto
STATE: Ontario
COUNTRY: Canada
ZIP: M5G 1R7
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/867,941
FILING DATE: 03-JUN-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Stewart, Michael I
REGISTRATION NUMBER: 24,973
REFERENCE/DOCKET NUMBER: 1038-681 MIS:jb
TELECOMMUNICATION INFORMATION:
TELEPHONE: (416) 595-1155
TELEFAX: (416) 595-1163
INFORMATION FOR SEQ ID NO: 6:
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SEQUENCE CHARACTERISTICS:
LENGTH: 7641 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-867-941-6

Query Match 0.6%; Score 19; DB 2; Length 7641;
Best Local Similarity 100.0%; Pred. No. 7.6;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2y 811 tcttggtctcaagcaaa 829
3b 3656 tcttggtctcaagcaaa 3674

RESULT 4

US-09-045-186-1/c
Sequence 1, Application US/09045186
Patent No. 6087154

GENERAL INFORMATION:
APPLICANT: Baez, Melvyn
TITLE OF INVENTION: RHESUS NEUROPEPTIDE Y1 RECEPTOR
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Eli Lilly and Company
STREET: Lilly Corporate Center
CITY: Indianapolis
STATE: Indiana
COUNTRY: United States of America
ZIP: 46285

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Gaylo, Paul J.
REGISTRATION NUMBER: 36,808
REFERENCE/DOCKET NUMBER: P-11376
TELECOMMUNICATION INFORMATION:
TELEPHONE: (317) 276-0756
TELEFAX: (317) 276-3861
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1152 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1152
US-09-045-186-1

Query Match 0.6%; Score 18; DB 3; Length 1152;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2y 2802 aaagaagcttggaaaca 2819
3b 1091 AAAGAAGCTTGGAAACA 1074

RESULT 5

US-09-045-186-3/c

Sequence 3, Application US/09045186
Patent No. 6087154
GENERAL INFORMATION:
APPLICANT: Baez, Melvyn
TITLE OF INVENTION: RHESUS NEUROPEPTIDE Y1 RECEPTOR
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Eli Lilly and Company
STREET: Lilly Corporate Center
CITY: Indianapolis
STATE: Indiana
COUNTRY: United States of America
ZIP: 46285
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Gaylo, Paul J.
REGISTRATION NUMBER: 36,808
REFERENCE/DOCKET NUMBER: P-11376
TELECOMMUNICATION INFORMATION:
TELEPHONE: (317) 276-0756
TELEFAX: (317) 276-3861
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1152 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: mRNA
US-09-045-186-3

Query Match 0.6%; Score 18; DB 3; Length 1152;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2802 aaagaagcttggaaaca 2819
Db 1091 AAAGAAGCTTGGAAACA 1074

RESULT 6

US-08-878-546-9
Sequence 9, Application US/08878546
Patent No. 5952463

GENERAL INFORMATION:
APPLICANT: SHIBANO, YUJI
APPLICANT: KIKUCHI, NORIHISA
TITLE OF INVENTION: NOVEL PROTEINASE INHIBITOR AND
TITLE OF INVENTION: GENE ENCODING THE INHIBITOR
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: STEINBERG, RASKIN & DAVIDSON P.C.
STREET: 1140 AVENUE OF THE AMERICAS
CITY: NEW YORK
STATE: NY
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

2y 2802 aaagaagcttggaaaca 2819
3b 1091 AAAGAAGCTTGGAAACA 1074

APPLICATION NUMBER: US/08/878,546
 FILING DATE: 19-JUN-1997
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: JP 158677/1996
 FILING DATE: 19-JUN-1996
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: JP 224104/1996
 FILING DATE: 26-AUG-1996
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: JP 48101/1997
 FILING DATE: 03-MAR-1997
 ATTORNEY/AGENT INFORMATION:
 NAME: DAVIDSON, CLIFFORD M.
 REGISTRATION NUMBER: 32,728
 REFERENCE/DOCKET NUMBER: 382.1009
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212)-768-3800
 TELEFAX: (212)382-2124
 INFORMATION FOR SEQ ID NO: 9:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 2186 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: double
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 ORIGINAL SOURCE:
 ORGANISM: STREPTOMYCES PLATENSIS
 STRAIN: Q268
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 1477..1911
 US-08-878-546-9

Query Match 0.6%; Score 18; DB 2; Length 2186;
 Best Local Similarity 100.0%; Pred.No. 23;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1652 ggtcctgggacccctggc 1669
 ||||||||||||||||
 S 1518 GGTCTGGGACCCCTGGC 1535

RESULT 7
 US-08-680-395-4/c
 Sequence 4, Application US/08680395
 Patent No. 5852010
 GENERAL INFORMATION:
 APPLICANT: Gray, Joe W.
 APPLICANT: Collins, Colin
 APPLICANT: Hwang, Soo-in
 APPLICANT: Godfrey, Tony
 APPLICANT: Kowbel, David
 APPLICANT: Rommens, Johanna
 TITLE OF INVENTION: Genes from the 20q13 Amplicon and Their
 NUMBER OF SEQUENCES: 40
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Townsend and Townsend and Crew LLP
 STREET: Two Embarcadero Center, Eighth Floor
 CITY: San Francisco
 STATE: California
 COUNTRY: USA
 ZIP: 94111-3834
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/680,395
 FILING DATE: 15-JUL-1996
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Bastian, Kevin L.
 REGISTRATION NUMBER: 34,774
 REFERENCE/DOCKET NUMBER: 023070-068900US
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (415) 576-0200
 TELEFAX: (415) 576-0300
 INFORMATION FOR SEQ ID NO: 4:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 2605 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: cDNA
 FEATURE:
 NAME/KEY: -
 LOCATION: 1..2605
 OTHER INFORMATION: /note= "cDNA clone cc43 of 4 kb
 OTHER INFORMATION: transcript"
 US-08-680-395-4

Query Match 0.6%; Score 18; DB 2; Length 2605;
 Best Local Similarity 100.0%; Pred.No. 23;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 448 ttccaaagtgtgtacttt 465
 ||||||||||||||||
 Db 1421 TTCCAAAGTGTGTACTTT 1404

RESULT 8
 US-08-967-101-23/c
 Sequence 23, Application US/08967101
 Patent No 5840540
 GENERAL INFORMATION:
 APPLICANT: ST. GEORGE-HYSLOP, PETER H
 APPLICANT: ROMMENS, JOHANNA M
 APPLICANT: FRASER, PAUL E
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 TO ALZHEIMER'S DISEASE
 NUMBER OF SEQUENCES: 193
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: TESTA, HURWITZ & THIBEAULT
 STREET: High Street Tower - 125 High Street
 CITY: Boston
 STATE: Massachusetts
 COUNTRY: U.S.A.
 ZIP: 02110
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/967,101
 FILING DATE: 10-NOV-1997
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/592,541
 FILING DATE:
 ATTORNEY/AGENT INFORMATION:
 NAME: Pitcher, Edmund R.
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (617) 248-7000
 TELEFAX: (617) 248-7100
 INFORMATION FOR SEQ ID NO: 23:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 289 base pairs
 TYPE: nucleic acid

STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
S-08-967-101-23

Query Match 0.6%; Score 17; DB 2; Length 289;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 2131 aagaggaagcagtgaa 2147
|||||
b 267 AAGAGGAAGCAGTGAA 251

RESULT 9

S-08-592-541-23/c
Sequence 23, Application US/08592541
Patent No. 5986054

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston

STATE: Massachusetts

COUNTRY: U.S.A.

ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
FILING DATE:

CLASSIFICATION: 800

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 23:

SEQUENCE CHARACTERISTICS:

LENGTH: 289 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

S-08-592-541-23

Query Match 0.6%; Score 17; DB 2; Length 289;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 2131 aagaggaagcagtgaa 2147
|||||
b 267 AAGAGGAAGCAGTGAA 251

RESULT 10

S-09-124-698-23/c
Sequence 23, Application US/09124698
Patent No. 611978

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H

APPLICANT: ROMMENS, JOHANNA M

APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston

STATE: Massachusetts

COUNTRY: U.S.A.

ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/592,541

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.

TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 23:

SEQUENCE CHARACTERISTICS:

LENGTH: 289 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-09-124-698-23

Query Match 0.6%; Score 17; DB 3; Length 289;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2131 aagaggaagcagtgaa 2147

Db 267 AAGAGGAAGCAGTGAA 251

RESULT 11

US-08-611-757-20/c
Sequence 20, Application US/08611757
Patent No. 5859230

GENERAL INFORMATION:

APPLICANT: Kim, Jungsuh P.

APPLICANT: Reyes, Gregory R.

APPLICANT: Wages, John

APPLICANT: Zhang-Keck, Zhen-Yang

APPLICANT: Young, Lavonne

TITLE OF INVENTION: NO. 5859230-A/No. 5859230-B/No. 5859230-C/No. 5859230-D/No.

NUMBER OF SEQUENCES: 106

CORRESPONDENCE ADDRESS:

ADDRESSEE: Dehlinger & Associates

STREET: 350 Cambridge Avenue, Suite 250

CITY: Palo Alto

STATE: CA

COUNTRY: USA

ZIP: 94306

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/611,757
 FILING DATE: 20-MAY-1994
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/246,985
 FILING DATE: 20-MAY-1994
 APPLICATION NUMBER: US 025,396
 FILING DATE: 24-FEB-1993
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 07/922,493
 FILING DATE: 30-JUL-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: Fabian, Gary R.
 REGISTRATION NUMBER: 33,875
 REFERENCE/DOCKET NUMBER: 4600-0201
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (415) 324-0880
 TELEFAX: (415) 324-0960
 INFORMATION FOR SEQ ID NO: 20:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 304 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: double
 TOPOLOGY: linear
 MOLECULE TYPE: DNA
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 ORIGINAL SOURCE:
 INDIVIDUAL ISOLATE: My 190 Clone D30
 PCT-08-611-757-20

Query Match 0.6%; Score 17; DB 2; Length 304;
 Best Local Similarity 100.0%; Pred. No. 71;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 132 cgcgcgagcgccgcg 148
 |||||
 Db 22 CGCCGCGAGCGCGCG 6

RESULT 12
 PCT-US95-05980-20/c
 Sequence 20, Application PC/TUS9505980
 GENERAL INFORMATION:
 APPLICANT:
 APPLICANT: Dehlinger & Associates
 TITLE OF INVENTION: Non-A/Non-B/Non-C/Non-D/Non-E Hepatitis
 TITLE OF INVENTION: Agents and Molecular Cloning Thereof
 NUMBER OF SEQUENCES: 106
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Dehlinger & Associates
 STREET: 350 Cambridge Avenue, Suite 250
 CITY: Palo Alto
 STATE: CA
 COUNTRY: USA
 ZIP: 94306
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: PCT/US95/05980
 FILING DATE:
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/246,986
 FILING DATE: 20-MAY-1994
 ATTORNEY/AGENT INFORMATION:
 NAME: Fabian, Gary R.
 REGISTRATION NUMBER: 33,875
 REFERENCE/DOCKET NUMBER: 4600-0201.49

TELECOMMUNICATION INFORMATION:
 TELEPHONE: (415) 324-0880
 TELEFAX: (415) 324-0960
 INFORMATION FOR SEQ ID NO: 20:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 304 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: double
 TOPOLOGY: linear
 MOLECULE TYPE: DNA
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 ORIGINAL SOURCE:
 INDIVIDUAL ISOLATE: My 190 Clone D30
 PCT-US95-05980-20
 Query Match 0.6%; Score 17; DB 4; Length 304;
 Best Local Similarity 100.0%; Pred. No. 71;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 132 cgcgcgagcgccgcg 148
 |||||
 Db 22 CGCCGCGAGCGCGCG 6

RESULT 13
 US-08-184-009-110
 Sequence 110, Application US/08184009
 Patent No. 5833975
 GENERAL INFORMATION:
 APPLICANT: Paoletti, Enzo
 APPLICANT: Tartaglia, James
 APPLICANT: Cox, William I.
 TITLE OF INVENTION: RECOMBINANT VIRUS IMMUNOTHERAPY
 NUMBER OF SEQUENCES: 217
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Curtis, Morris & Safford
 STREET: 530 Fifth Avenue
 CITY: New York
 STATE: NY
 COUNTRY: USA
 ZIP: 10036
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/184,009
 FILING DATE: 19-JAN-1994
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Frommer, William S.
 REGISTRATION NUMBER: 25,506
 REFERENCE/DOCKET NUMBER: 454310-2530
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212) 840-3333
 TELEFAX: (212) 840-0712
 TELEX: 425066CURTMS
 INFORMATION FOR SEQ ID NO: 110:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1084 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: CDNA
 US-08-184-009-110

Query Match 0.6%; Score 17; DB 2; Length 1084;
 Best Local Similarity 100.0%; Pred. No. 72;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

REGISTRATION NUMBER: 25,506
REFERENCE/DOCKET NUMBER: 454310-2530
TELEPHONE: (212) 840-3333
TELEFAX: (212) 840-0712
TELEX: 425066CURTMS
INFORMATION FOR SEQ ID NO: 109:
SEQUENCE CHARACTERISTICS:
LENGTH: 1094 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-458-356-109

Query Match 0.6%; Score 17; DB 2; Length 1094;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1652 ggtctctgggcaccctgg 1668
Db 314 GGTCTCTGGGCACCCCTGG 330

RESULT 17
US-08-993-118-8
; Sequence 8, Application US/08993118
; Patent No. 5997872
; GENERAL INFORMATION:
; APPLICANT: LUCAS, Sophie;
; APPLICANT: DE SMET, Charles;
; APPLICANT: BOON-FALLEUR, Thierry
; TITLE OF INVENTION: ISOLATED NUCLEIC ACID MOLECULE CODING FOR TUMOR
; TITLE OF INVENTION: REJECTION ANTIGEN PRECURSOR MAGE-C1 AND USES
; TITLE OF INVENTION: THEREOF
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Felfe & Lynch
; STREET: 805 Third Avenue
; CITY: New York City
; STATE: New York
; COUNTRY: USA
; ZIP: 10022
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: PC-DOS
; SOFTWARE: Wordperfect
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/993,118
; FILING DATE: April 25, 1997
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/845,528
; FILING DATE: April 25, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Mary Anne Schofield
; REGISTRATION NUMBER: 36,669
; REFERENCE/DOCKET NUMBER: LUD 5455
; TELEPHONE: (212) 688-9200
; TELEFAX: (212) 838-3884
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1691 base pairs
; TYPE: nucleotides
; STRANDEDNESS: single stranded
; TOPOLOGY: linear
US-08-993-118-8

Query Match 0.6%; Score 17; DB 2; Length 1691;

Best Local Similarity 100.0%; Pred. No. 72;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1652 ggtctctgggcaccctgg 1668
Db 317 GGTCTCTGGGCACCCCTGG 333

RESULT 18
US-08-845-528C-8
; Sequence 8, Application US/08845528C
; Patent No. 6027924
; GENERAL INFORMATION:
; APPLICANT: LUCAS, Sophie;
; APPLICANT: DE SMET, Charles;
; APPLICANT: BOON-FALLEUR, Thierry
; TITLE OF INVENTION: ISOLATED NUCLEIC ACID MOLECULE CODING FOR TUMOR
; TITLE OF INVENTION: REJECTION ANTIGEN PRECURSOR MAGE-C1 AND USES
; TITLE OF INVENTION: THEREOF
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Felfe & Lynch
; STREET: 805 Third Avenue
; CITY: New York City
; STATE: New York
; COUNTRY: USA
; ZIP: 10022

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
COMPUTER: IBM PS/2
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/845,528C
FILING DATE: April 25, 1997
CLASSIFICATION: 4335
ATTORNEY/AGENT INFORMATION:
NAME: Mary Anne Schofield
REGISTRATION NUMBER: 36,669
REFERENCE/DOCKET NUMBER: LUD 5455
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 1691 base pairs
TYPE: nucleotides
STRANDEDNESS: single stranded
TOPOLOGY: linear
US-08-845-528C-8

Query Match 0.6%; Score 17; DB 3; Length 1691;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1652 ggtctctgggcaccctgg 1668
Db 317 GGTCTCTGGGCACCCCTGG 333

RESULT 19
US-08-951-148-2
; Sequence 2, Application US/08951148
; Patent No. 5871973
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Bandman, Olga
; APPLICANT: Kal, Preeti
; APPLICANT: Shah, Purvi
; TITLE OF INVENTION: CELL DIVISION REGULATORS
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:

ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/951,148
FILING DATE: Herewith
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0407 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1816 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: SPLNFT01
CLONE: 26459

Query Match 0.6%; Score 17; DB 2; Length 1816;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 804 ggaacctcttggtgct 820
|||||

Db 717 GGAACTCTTGCTGCT 733

RESULT 20

US-09-165-234-2
Sequence 2, Application US/09165234
Patent No. 5928899
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
APPLICANT: Lal, Preeti
APPLICANT: Shah, Purvi
TITLE OF INVENTION: CELL DIVISION REGULATORS
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/165,234
FILING DATE:

CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/951,148
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0407 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1816 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: SPLNFT01
CLONE: 26459

US-09-165-234-2

Query Match 0.6%; Score 17; DB 2; Length 1816;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 804 ggaacctcttggtgct 820
|||||

Db 717 GGAACTCTTGCTGCT 733

RESULT 21

US-09-274-570-2
Sequence 2, Application US/09274570
Patent No. 6121019
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
APPLICANT: Lal, Preeti
APPLICANT: Shah, Purvi
TITLE OF INVENTION: CELL DIVISION REGULATORS
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/274,570
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/951,148
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0407 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:

LENGTH: 1816 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 IMMEDIATE SOURCE:
 LIBRARY: SPLNFZT01
 CLONE: 26459
 S-09-274-570-2

Query Match 0.6%; Score 17; DB 3; Length 1816;
 Best Local Similarity 100.0%; Pred. No. 72;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 804 ggaactctttgtgct 820
 |||||
 DB 717 GGAACCTCTGGTCT 733

RESULT 22

US-07-807-043B-7
 ; Sequence 7, Application US/07807043B
 ; Patent No. 5342774

GENERAL INFORMATION:

APPLICANT: Boon, Thierry, Van den Eynde, Beno t
 TITLE OF INVENTION: Tumor Rejection Antigen Precursors, Tumor
 TITLE OF INVENTION: Rejection Antigens and Uses Thereof
 NUMBER OF SEQUENCES: 16
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Felfe & Lynch
 STREET: 805 Third Avenue
 CITY: New York City
 STATE: New York
 ZIP: 10022

COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage

COMPUTER: IBM
 OPERATING SYSTEM: PC-DOS

SOFTWARE: Wordperfect
 CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/07/807,043B
 FILING DATE: 19911212

CLASSIFICATION: 424

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/764,364
 FILING DATE: 23-SEPTEMBER-1991

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/728,838
 FILING DATE: 9-JULY-1991

PRIOR APPLICATION DATA: 07/705,702
 FILING DATE: 23-May-1991

ATTORNEY/AGENT INFORMATION:

NAME: Hanson, No. 5342774man D.
 REGISTRATION NUMBER: 30,946

REFERENCE/DOCKET NUMBER: LUD 253.3
 TELEPHONE: (212) 688-9200

TELEFAX: (212) 838-3884

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 2419 base pairs

TYPE: NUCLEIC ACID

STRANDEDNESS: singular

TOPOLOGY: linear

MOLECULE TYPE: genomic DNA

US-07-807-043B-7

Query Match 0.6%; Score 17; DB 1; Length 2419;
 Best Local Similarity 100.0%; Pred. No. 73;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggacccctgg 1668
 |||||
 DB 739 GGTCTGGGACCCCTGG 755

RESULT 23

US-08-299-849B-7
 ; Sequence 7, Application US/08299849B
 ; Patent No. 5612201

GENERAL INFORMATION:

APPLICANT: De Plaeen, Etienne; Boon-Falleur, Thierry;
 APPLICANT: Leth, Bernard; Szikora, Jean-Pierre; De Smet, Charles;
 APPLICANT: Chomez, Patrick
 TITLE OF INVENTION: Isolated Nucleic Acid Molecules Useful In
 TITLE OF INVENTION: Determining Expression Of A Tumor Antigen Precursor.
 NUMBER OF SEQUENCES: 48
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Felfe & Lynch
 STREET: 805 Third Avenue
 CITY: New York City
 STATE: New York
 ZIP: 10022

COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage

COMPUTER: IBM
 OPERATING SYSTEM: PC-DOS

SOFTWARE: Wordperfect
 CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/299,849B
 FILING DATE: 1-SEPTEMBER-1994

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/037,230
 FILING DATE: 26-MARCH-1993

PRIOR APPLICATION DATA:

APPLICATION NUMBER: PCT/US92/04354
 FILING DATE: 22-MAY-1992

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/807,043
 FILING DATE: 12-DECEMBER-1991

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/764,364
 FILING DATE: 23-SEPTEMBER-1991

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/728,838
 FILING DATE: 9-JULY-1991

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 07/705,702
 FILING DATE: 23-May-1991

ATTORNEY/AGENT INFORMATION:

NAME: Hanson, No. 5612201man D.
 REGISTRATION NUMBER: 30,946

REFERENCE/DOCKET NUMBER: LUD 5355
 TELEPHONE: (212) 688-9200

TELEFAX: (212) 838-3884

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 2419 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: genomic DNA

US-08-299-849B-7

Query Match 0.6%; Score 17; DB 1; Length 2419;
 Best Local Similarity 100.0%; Pred. No. 73;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggacccctgg 1668
 |||||
 DB 739 GGTCTGGGACCCCTGG 755

RESULT 24
US-08-142-368A-7
Sequence 7, Application US/08142368A
Patent No. 5925729
GENERAL INFORMATION:
APPLICANT: Boon-Falleur, Thierry; Van der Bruggen, Thierry;
APPLICANT: Van den Eynde, Beno t.; Van Pel, Aline; De Plaen, Etienne;
APPLICANT: Lurquin, Christophe; Chomez, Patrick; Traversari, Catia
TITLE OF INVENTION: Tumor Rejection Antigen Precursors, Tumor
TITLE OF INVENTION: Rejection Antigens and Uses Thereof
NUMBER OF SEQUENCES: 26
CORRESPONDENCE ADDRESS:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/142,368A
FILING DATE: 02-MAY-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/04354
FILING DATE: 22-MAY-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/807,043
FILING DATE: 12-DECEMBER-1991
APPLICATION NUMBER: 07/764,364
FILING DATE: 23-SEPTEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
FILING DATE: 9-JULY-1991
APPLICATION NUMBER: 07/705,702
FILING DATE: 23-MAY-1991
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 5925729man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5253.4-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2419 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
US-08-142-368A-7
Query Match 0.6%; Score 17; DB 2; Length 2419;
Best Local Similarity 100.0%; Pred.No.73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CY 1652 ggtcctgggaccctgg 1668
|||||
DB 739 GGTCTGGGACCCCTGG.755
RESULT 25
US-08-967-727-7
Sequence 7, Application US/08967727
Patent No. 6025474

GENERAL INFORMATION:
APPLICANT: Gaugler, B atrice; Van den Eynde, Beno t.;
APPLICANT: van der Bruggen, Pierre; Boon-Falleur, Thierry
TITLE OF INVENTION: Isolated Nucleic Acid Molecules Coding For
TITLE OF INVENTION: Tumor Rejection Antigen Precursor Wage-3 And Uses Thereof
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,727
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/037,230
FILING DATE: 26-MARCH-1993
APPLICATION NUMBER: PCT/US92/04354
FILING DATE: 22-MAY-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/807,043
FILING DATE: 12-DECEMBER-1991
APPLICATION NUMBER: 07/764,365
FILING DATE: 23-SEPTEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
FILING DATE: 9-JULY-1991
APPLICATION NUMBER: 07/705,702
FILING DATE: 23-MAY-1991
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 6025474man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5353
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2419 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
US-08-967-727-7
Query Match 0.6%; Score 17; DB 3; Length 2419;
Best Local Similarity 100.0%; Pred.No.73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CY 1652 ggtcctgggaccctgg 1668
|||||
DB 739 GGTCTGGGACCCCTGG 755
RESULT 26
US-08-465-167A-23-
Sequence 23, Application US/08465167A
Patent No. 5750395
GENERAL INFORMATION:
APPLICANT: Fikes, John D.
APPLICANT: Livingston, Brian D.
APPLICANT: Sette, Alessandro D.
APPLICANT: Sidney, John C.

TITLE OF INVENTION: DNA ENCODING MAGE-1 C-TERMINAL
TITLE OF INVENTION: IMMUNOGENIC PEPTIDES (as amended)
NUMBER OF SEQUENCES: 51
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, 8th Floor
CITY: San Francisco
STATE: CA
COUNTRY: USA
ZIP: 94111
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/465,167A
FILING DATE: 05-JUN-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/103,623
FILING DATE: 06-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: Parmelee, Steven W.
REGISTRATION NUMBER: 31,990
REFERENCE/DOCKET NUMBER: 14137-60-1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 206-467-9600
TELEFAX: 415-576-0300
INFORMATION FOR SEQ ID NO: 23:
SEQUENCE CHARACTERISTICS:
LENGTH: 2420 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
FEATURE:
NAME/KEY: CDS
LOCATION: 626..1552
US-08-465-167A-23

Query Match 0.6%; Score 17; DB 1; Length 2420;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctggggcaccctgg 1668
|||||
Q 739 GGTCTGGGACCCCTGG 755

RESULT 27
US-08-295-882-1/C
Sequence 1, Application US/08295882
Patent No. 5569833
GENERAL INFORMATION:
APPLICANT:
TITLE OF INVENTION: METHOD FOR ENHANCING
TITLE OF INVENTION: PLANT PRECOCITY AND/OR REDUCING THE
TITLE OF INVENTION: STORED NITRATE CONTENT OF A PLANT
NUMBER OF SEQUENCES: 1
CORRESPONDENCE ADDRESS:
ADDRESSEE: Christle, Parker & Hale
STREET: P.O. Box 7068
CITY: Pasadena
STATE: CA
COUNTRY: USA
ZIP: 91109-7068
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: WORD PERFECT 5.1 release

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/295,882
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/FR 93/00222
FILING DATE: March 5, 1993
APPLICATION NUMBER: 92 02658
FILING DATE: March 5, 1992
ATTORNEY/AGENT INFORMATION:
NAME: Prout, D. Bruce
REGISTRATION NUMBER: 20,958
REFERENCE/DOCKET NUMBER: 27209/DBP
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 3457 base pairs
TYPE: nucleotide with corresponding
TYPE: protein
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
ORIGINAL SOURCE:
ORIGINAL SOURCE: (1.VI.A) ORGANISM: Nicotiana tabacum
ORIGINAL SOURCE: (1.VI.B) CELL LINE: N. tabacum cv. Xanthi
IMMEDIATE SOURCE: Leaf
FEATURE:
NAME/KEY: Nitrate reductase
LOCATION: from 1 to 143 bp: Leader
LOCATION: non translated 5 sequence (leader)
LOCATION: from 144 to 2855 bp: coding sequence
LOCATION: for nitrate reductase apoenzyme
LOCATION: from 2856 to 3457 bp: non translated
LOCATION: 3 sequence
US-08-295-882-1
Query Match 0.6%; Score 17; DB 1; Length 3457;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 909 gaaggagagagatttt 925
|||||
Db 94 GAAGGAAGAGAGATT TT 78
RESULT 28
US-08-343-760A-1
Sequence 1, Application US/08343760A
Patent No. 5679783
GENERAL INFORMATION:
APPLICANT: De Robertis, Edward M
APPLICANT: Sasal, Yoshiki
TITLE OF INVENTION: Tissue Differentiation Affecting
TITLE OF INVENTION: Factor and Composition
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Majestic, Parsons, Siebert & Hsue
STREET: Four Embarcadero Center, Suite 1450
CITY: San Francisco
STATE: CA
COUNTRY: USA
ZIP: 94111
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/343,760A
FILING DATE: 22-NOV-1994
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Siebert, J. Suzanne

REGISTRATION NUMBER: 28,758
REFERENCE/DOCKET NUMBER: 3100.1
TELEPHONE: (415) 363-5556
TELEFAX: (415) 362-5418
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 3796 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-08-343-760A-1

Query Match 0.6%; Score 17; DB 1; Length 3796;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Y 270 ctctacgtctctccga 286
|||||
Db 1036 CTCACGCTCTCCGA 1052

RESULT 29
US-08-441-430-1
Sequence 1, Application US/08441430
Patent No. 5681942
GENERAL INFORMATION:
APPLICANT: Buchwald, Manuel
APPLICANT: Strathdee, Craig A.
APPLICANT: Wevrick, Rachel
APPLICANT: Mathew, Christopher George Porter
TITLE OF INVENTION: Fanconi Anemia Type C Gene
NUMBER OF SEQUENCES: 73
CORRESPONDENCE ADDRESS:
ADDRESSEE: Richard J. Polley, Esq.
ADDRESSEE: Klarquist, Sparkman, Campbell, Leigh &
ADDRESSEE: Winston, LLP
STREET: 121 S.W. Salmon, Suite 1600
CITY: Portland
STATE: Oregon
COUNTRY: U.S.A.
ZIP: 97204
COMPUTER READABLE FORM:
MEDIUM TYPE: Disk, 3+-inch
COMPUTER: IBM PC compatible
OPERATING SYSTEM: MS DOS
SOFTWARE: WordPerfect 5.1/ASCII Text File
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/441.430
FILING DATE: May 15, 1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: U.S. 07/876,285
FILING DATE: April 29, 1992
APPLICATION NUMBER: U.S. 07/918,313
FILING DATE: July 21, 1992
APPLICATION NUMBER: U.S. 08/003,963
FILING DATE: January 15, 1993
ATTORNEY/AGENT INFORMATION:
NAME: Richard J. Polley, Esq.
REGISTRATION NUMBER: 28,107
REFERENCE/DOCKET NUMBER: 3812-42824
TELEPHONE: (503) 226-7391
TELEFAX: (503) 228-9446
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 4488 base pairs
TYPE: Nucleic Acid
STRANDEDNESS: Double stranded
TOPOLOGY: Linear

MOLECULE TYPE: CDNA to mRNA
HYPOTHEICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Homo sapiens
IMMEDIATE SOURCE:
LIBRARY: Human cDNA
POSITION IN GENOME: (of corresponding genomic gene)
CHROMOSOME/SEGMENT: 9q
MAP POSITION: 22.3
UNITS:
US-08-441-430-1
Query Match 0.6%; Score 17; DB 1; Length 4488;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2105 tgaagccaccctggaag 2121
|||||
Db 4189 TGAAGCCACCCTGGAAG 4205
RESULT 30
US-07-807-043B-8
Sequence 8, Application US/07807043B
Patent No. 5342774
GENERAL INFORMATION:
APPLICANT: Boon, Thierry, Van den Eynde, Beno t
TITLE OF INVENTION: Tumor Rejection Antigen Precursors, Tumor
TITLE OF INVENTION: Rejection Antigens and Uses Thereof
NUMBER OF SEQUENCES: 16
CORRESPONDENCE ADDRESS:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/807,043B
FILING DATE: 19911212
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/764,364
FILING DATE: 23-SEPTEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
FILING DATE: 9-JULY-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/705,702
FILING DATE: 23-May-1991
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 5342774man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 253.3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 5674 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: singular
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
FEATURE:
NAME/KEY: MAGE-1 gene
US-07-807-043B-8

Query Match 0.6%; Score 17; DB 1; Length 5674;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggaccctgg 1668
|||||
DB 3994 GGTCTGGGACCCCTGG 4010

RESULT 31

US-08-190-411A-1
; Sequence 1, Application US/08190411A
; Patent No. 5541104
; GENERAL INFORMATION:
; APPLICANT: Chen, Yao-Tsung; Stockert, Elisabeth;
; APPLICANT: Chen, Yachi; Garin-Chesa, Pilar; Rettig, Wolfgang J.;
; APPLICANT: van der Bruggen, Pierre; Boon-Falleur, Thierry;
; APPLICANT: Old, Lloyd J.
; TITLE OF INVENTION: MONOCLONAL ANTIBODIES WHICH BIND TO
; TITLE OF INVENTION: TUMOR REJECTION ANTIGEN PRECURSOR MAGE-1, RECOMBINANT MAGE-1,
; TITLE OF INVENTION: AND MAGE-1 DERIVED IMMUNOGENIC PEPTIDES
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Felfe & Lynch
; STREET: 805 Third Avenue
; CITY: New York City
; STATE: New York
; ZIP: 10022
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
; COMPUTER: IBM
; OPERATING SYSTEM: PC-DOS
; SOFTWARE: Wordperfect
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/190,411A
; FILING DATE: 01-FEBRUARY-1994
; CLASSIFICATION: 436
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 037,230
; FILING DATE: 26-MARCH-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/04354
; FILING DATE: 22-MAY-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/807,043
; FILING DATE: 12-DECEMBER-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/764,364
; FILING DATE: 23-SEPTEMBER-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/728,838
; APPLICATION NUMBER: 9-JULY-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/705,702
; FILING DATE: 23-MAY-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Hanson, No. 5541104man D.
; REGISTRATION NUMBER: 30,946
; REFERENCE/DOCKET NUMBER: LUD 5354
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 688-9200
; TELEFAX: (212) 838-3884
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 5674 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
; FEATURE:
; NAME/KEY: MAGE-1 gene

US-08-190-411A-1

Query Match 0.6%; Score 17; DB 1; Length 5674;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggaccctgg 1668
|||||
DB 3994 GGTCTGGGACCCCTGG 4010

RESULT 32

US-08-299-849B-8
; Sequence 8, Application US/08299849B
; Patent No. 5612201
; GENERAL INFORMATION:
; APPLICANT: De Plaen, Etienne; Boon-Falleur, Thierry;
; APPLICANT: Leth, Bernard; Szikora, Jean-Pierre; De Smet, Charles;
; APPLICANT: Chomez, Patrick
; TITLE OF INVENTION: Isolated Nucleic Acid Molecules Useful In
; TITLE OF INVENTION: Determining Expression Of A Tumor Antigen Precursor
; NUMBER OF SEQUENCES: 48
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Felfe & Lynch
; STREET: 805 Third Avenue
; CITY: New York City
; STATE: New York
; ZIP: 10022
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
; COMPUTER: IBM
; OPERATING SYSTEM: PC-DOS
; SOFTWARE: Wordperfect
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/299,849B
; FILING DATE: 1-SEPTEMBER-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/037,230
; FILING DATE: 26-MARCH-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/04354
; FILING DATE: 22-MAY-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/807,043
; FILING DATE: 12-DECEMBER-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/764,364
; FILING DATE: 23-SEPTEMBER-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/728,838
; APPLICATION NUMBER: 9-JULY-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/705,702
; FILING DATE: 23-MAY-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Hanson, No. 5612201man D.
; REGISTRATION NUMBER: 30,946
; REFERENCE/DOCKET NUMBER: LUD 5355
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 688-9200
; TELEFAX: (212) 838-3884
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 5674 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
; FEATURE:
; NAME/KEY: MAGE-1 gene

US-08-299-849B-8

Query Match 0.6%; Score 17; DB 1; Length 5674;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1652 ggtcctgggacccctgg 1668
|||||
Db 3994 GGTCTGGGACCCCTGG 4010

RESULT 33
US-08-560-024-1
Sequence 1, Application US/08560024
Patent No. 5843448
GENERAL INFORMATION:
APPLICANT: Chen, Yao-Tseng; Stockert, Elisabeth;
APPLICANT: Chen, Yachi; Garin-Chesa, Pilar; Rettig, Wolfgang J.;
APPLICANT: van der Bruggen, Pierre; Boon-Falleur, Thierry;
APPLICANT: Old, Lloyd J.
TITLE OF INVENTION: MONOCLONAL ANTIBODIES WHICH BIND TO
TITLE OF INVENTION: TUMOR REJECTION ANTIGEN PRECURSOR MAGE-1, RECOMBINANT MAGE-1,
TITLE OF INVENTION: AND MAGE-1 DERIVED IMMUNOGENIC PEPTIDES
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/560,024
FILING DATE:
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/190,411
FILING DATE: 01-FEBRUARY-1994
APPLICATION NUMBER: 037,230
FILING DATE: 26-MARCH-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/04354
FILING DATE: 22-MAY-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/807,043
FILING DATE: 12-DECEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/764,364
FILING DATE: 23-SEPTEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
FILING DATE: 9-JULY-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/705,702
FILING DATE: 23-MAY-1991
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 5843448man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5354
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5674 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA

FEATURE:
NAME/KEY: MAGE-1 gene
US-08-560-024-1

Query Match 0.6%; Score 17; DB 2; Length 5674;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1652 ggtcctgggacccctgg 1668
|||||
Db 3994 GGTCTGGGACCCCTGG 4010

RESULT 34
US-08-142-368A-8
Sequence 8, Application US/08142368A
Patent No. 5925729
GENERAL INFORMATION:
APPLICANT: Boon-Falleur, Thierry; Van der Bruggen, Thierry;
APPLICANT: Van den Eynde, Beno t; Van Pel, Aline; De Plaen, Etienne;
APPLICANT: Lurquin, Christophe; Chomez, Patrick; Traversari, Catia
TITLE OF INVENTION: Tumor Rejection Antigen Precursors, Tumor
TITLE OF INVENTION: Rejection Antigens and Uses Thereof
NUMBER OF SEQUENCES: 26
CORRESPONDENCE ADDRESS:
ADDRESSEE: Felfe & Lynch
STREET: 805 Third Avenue
CITY: New York City
STATE: New York
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
COMPUTER: IBM
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/142,368A
FILING DATE: 02-MAY-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US92/04354
FILING DATE: 22-MAY-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/807,043
FILING DATE: 12-DECEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/764,364
FILING DATE: 23-SEPTEMBER-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/728,838
FILING DATE: 9-JULY-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/705,702
FILING DATE: 23-MAY-1991
ATTORNEY/AGENT INFORMATION:
NAME: Hanson, No. 5925729man D.
REGISTRATION NUMBER: 30,946
REFERENCE/DOCKET NUMBER: LUD 5253.4-US
TELEPHONE: (212) 688-9200
TELEFAX: (212) 838-3884
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 5674 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
FEATURE:
NAME/KEY: MAGE-1 gene
US-08-142-368A-8

Query Match 0.6%; Score 17; DB 2; Length 5674;
 Best Local Similarity 100.0%; Pred. No. 73;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggacccctgg 1668
 |||||
 DB 3994 GGTCTGGGACCCCTGG 4010

RESULT 35
 US-08-967-727-8
 ; Sequence 8, Application US/08967727
 ; Patent No. 6025474
 ; GENERAL INFORMATION:
 ; APPLICANT: Gaugler, B atrice; Van den Eynde, Beno t;
 ; APPLICANT: van der Bruggen, Pierre; Boon-Falleur, Thierry
 ; TITLE OF INVENTION: Isolated Nucleic Acid Molecules Coding For
 ; TITLE OF INVENTION: Tumor Rejection Antigen Precursor Mage-3 And Uses Thereof
 ; NUMBER OF SEQUENCES: 30
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Felife & Lynch
 ; STREET: 805 Third Avenue
 ; CITY: New York City
 ; STATE: New York
 ; ZIP: 10022
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Diskette, 5.25 inch, 360 kb storage
 ; COMPUTER: IBM
 ; OPERATING SYSTEM: PC-DOS
 ; SOFTWARE: Wordperfect
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/967,727
 ; FILING DATE:
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 08/037,230
 ; FILING DATE: 26-MARCH-1993
 ; APPLICATION NUMBER: PCT/US92/04354
 ; FILING DATE: 22-MAY-1992
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 07/807,043
 ; FILING DATE: 12-DECEMBER-1991
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 07/764,365
 ; FILING DATE: 23-SEPTEMBER-1991
 ; APPLICATION NUMBER: 07/728,838
 ; FILING DATE: 9-JULY-1991
 ; PRIOR APPLICATION DATA: 07/705,702
 ; FILING DATE: 23-MAY-1991
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Hanson, No. 6025474man D.
 ; REGISTRATION NUMBER: 30,946
 ; REFERENCE/DOCKET NUMBER: LUD 5353
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (212) 688-9200
 ; TELEFAX: (212) 838-3884
 ; INFORMATION FOR SEQ ID NO: 8:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 5674 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: genomic DNA
 ; FEATURE:
 ; NAME/KEY: MAGE-1 gene
 ; US-08-967-727-8

Query Match 0.6%; Score 17; DB 3; Length 5674;
 Best Local Similarity 100.0%; Pred. No. 73;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggacccctgg 1668
 |||||
 DB 3994 GGTCTGGGACCCCTGG 4010

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1652 ggtcctgggacccctgg 1668
 |||||
 DB 3994 GGTCTGGGACCCCTGG 4010

RESULT 36
 US-08-321-478-6
 ; Sequence 6, Application US/08321478
 ; Patent No. 5527677
 ; GENERAL INFORMATION:
 ; APPLICANT: DESUCHI, Takeo
 ; APPLICANT: KINOSHITA, Moritoshi
 ; APPLICANT: KATSURAGI, Kiyonori
 ; APPLICANT: SHIN, Sadahito
 ; TITLE OF INVENTION: HUMAN ARYLAMINE N-ACETYLTRANSFERASE
 ; TITLE OF INVENTION: GENES
 ; NUMBER OF SEQUENCES: 13
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Sughrue, Mion, Zinn, Macpeak & Seas
 ; STREET: 2100 Pennsylvania Avenue, N.W.
 ; CITY: Washington
 ; STATE: D.C.
 ; COUNTRY: United States
 ; ZIP: 20037-3202
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.25
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/321,478
 ; FILING DATE: 11-OCT-1994
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/038,667
 ; FILING DATE: 23-MAR-1993
 ; APPLICATION NUMBER: JP 64669/1992
 ; FILING DATE: 23-MAR-1992
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (202) 293-7060
 ; TELEFAX: (202) 293-7860
 ; TELEX: 6491103
 ; INFORMATION FOR SEQ ID NO: 6:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 6464 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: DNA (genomic)
 ; FEATURE:
 ; NAME/KEY: CDS
 ; LOCATION: 723..1595
 ; FEATURE:
 ; NAME/KEY: exon
 ; LOCATION: 717..1936
 ; FEATURE:
 ; NAME/KEY: polyA_signal
 ; LOCATION: 1794..1799
 ; FEATURE:
 ; NAME/KEY: polyA_signal
 ; LOCATION: 1800..1805
 ; US-08-321-478-6

Query Match 0.6%; Score 17; DB 1; Length 6464;
 Best Local Similarity 100.0%; Pred. No. 74;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1032 aggtaccacgaaggaagc 1048
 |||||
 DB 5916 AGGTACCAAGGAAGGC 5932


```
RESULT 37
US-07-853-913-1
Sequence 1, Application US/07853913
Patent No. 5338839
GENERAL INFORMATION:
APPLICANT: McKay, Ronald D.G.
APPLICANT: Lendahl, Urban
TITLE OF INVENTION: Nestin Expression As An Indicator of
TITLE OF INVENTION: Neuroepithelial Tumors
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
STREET: Two Militia Drive
CITY: Lexington
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02173
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/853,913
FILING DATE: 19920319
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/660,412
FILING DATE: 22-FEB-1991
PRIOR APPLICATION DATA: US 07/603,803
FILING DATE: 25-OCT-1990
APPLICATION NUMBER: US 07/201,762
FILING DATE: 02-JUN-1988
APPLICATION NUMBER: US 07/180,548
ATTORNEY/AGENT INFORMATION:
NAME: Granahan, Patricia
REGISTRATION NUMBER: 32,227
REFERENCE/DOCKET NUMBER: MIT-4641AAAA
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-861-6240
TELEFAX: 617-861-9340
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 11236 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-07-853-913-1

Query Match 0.6%; Score 17; DB 1; Length 11236;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

-Y 1877 ccttcaggaggcgctg 1893
|||||
2b 3466 CCTTCAGGAGGGGCTG 3482

RESULT 38
US-08-781-891-207
Sequence 207, Application US/08781891
Patent No. 6090620
GENERAL INFORMATION:
APPLICANT: Fu, Ying-Hui
APPLICANT: Yu, Chang-En
APPLICANT: Oshima, Junko
```

```
APPLICANT: Mulligan, John T.
APPLICANT: Schellenberg, Gerald D.
TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO
TITLE OF INVENTION: WERNER'S SYNDROME
NUMBER OF SEQUENCES: 209
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104-7092
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/781,891
FILING DATE: 27-DEC-1996
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION:
NAME: No. 6090620tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 240052.419
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 207:
SEQUENCE CHARACTERISTICS:
LENGTH: 29604 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-781-891-207

Query Match 0.6%; Score 17; DB 3; Length 29604;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 525 aaaggaataagaactggc'541
|||||
DB 14826 AAAGGAATAGACTGGC 14842

RESULT 39
US-09-335-409-1/c
Sequence 1, Application US/09335409
Patent No. 6121029
GENERAL INFORMATION:
APPLICANT: Schupp, Thomas
APPLICANT: Ligon, James
APPLICANT: Molnar, Istvan
APPLICANT: Zirkle, Ross
APPLICANT: Cyr, Devon
APPLICANT: Goslach, Joern
TITLE OF INVENTION: GENES FOR THE BIOSYNTHESIS OF EPOTHILONES
FILE REFERENCE: 4-30582A
CURRENT APPLICATION NUMBER: US/09/335,409
CURRENT FILING DATE: 1999-06-17
NUMBER OF SEQ ID NOS: 30
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 1
LENGTH: 68750
TYPE: DNA
ORGANISM: Sorangium cellulosum
US-09-335-409-1

Query Match 0.6%; Score 17; DB 3; Length 68750;
Best Local Similarity 100.0%; Pred. No. 75;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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7 1457 cggccagcccccagcag 1473
8 |||||||||||||||
9 20332 CGGCCAGCCCCAGCAG 20316

RESULT 40
US-07-753-110B-12
: Sequence 12, Application US/07753110B
: Patent No. 5436141
: GENERAL INFORMATION:
: APPLICANT: Miyata, Shohei
: APPLICANT: Ohshima, Atsushi
: APPLICANT: Inouye, Sumiko
: APPLICANT: Inouye, Masayori
: TITLE OF INVENTION: METHOD FOR SYNTHESIZING STABLE
: TITLE OF INVENTION: SINGLE-STRANDED CDNA IN EUKARYOTES BY MEANS OF A BACTERIAL
: NUMBER OF SEQUENCES: 20
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Weiser & Associates
: STREET: 230 South Fifteenth Street, Suite 500
: CITY: Philadelphia
: STATE: Pennsylvania
: COUNTRY: U.S.A.
: ZIP: 19102
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: FILING DATE: 30-AUG-1991
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: Weiser, Gerard J.
: REGISTRATION NUMBER: 19,763
: REFERENCE/DOCKET NUMBER: 377.5584P
: TELEPHONE: 215-875-8383
: TELEFAX: 215-875-8394
: INFORMATION FOR SEQ ID NO: 12:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 76 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: 19
: OTHER INFORMATION: /note= "The 2' position of this
: OTHER INFORMATION: nucleotide is linked to the 5' position of
: OTHER INFORMATION: nucleotide number 1 of SEQ ID NO: 11 of this
: OTHER INFORMATION: application."
: FEATURE:
: NAME/KEY: misc_binding
: LOCATION: 69..76
: OTHER INFORMATION: /note= "This region can hydrogen
: OTHER INFORMATION: bond to nucleotides 156-163 of SEQ ID NO: 11 of
: OTHER INFORMATION: this application."
US-07-753-110B-12
: Query Match 0.5%; Score 16; DB 1; Length 76;
: Best Local Similarity 87.5%; Pred. No. 2.2e+02;
: Matches 14; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

2165 gtcccaagccatcagc 2180
1:|||||||||:|||||
5 GUCCCAAGCCCAUCCAGC 20
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```

RESULT 41
US-08-503-730-16
: Sequence 16, Application US/08503730
: Patent No. 5780269
: GENERAL INFORMATION:
: APPLICANT: Inouye, Sumiko
: APPLICANT: Inouye, Masayori
: TITLE OF INVENTION: NEW HYBRID MOLECULES
: NUMBER OF SEQUENCES: 45
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Weiser & Associates
: STREET: 230 South Fifteenth Street Suite 500
: CITY: Philadelphia
: STATE: PA
: COUNTRY: USA
: ZIP: 19102
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: FILING DATE: 18-JUL-1995
: CLASSIFICATION: 435
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/817,430
: FILING DATE: 06-JAN-1992
: ATTORNEY/AGENT INFORMATION:
: NAME: Weiser, Gerard J.
: REGISTRATION NUMBER: 19,763
: REFERENCE/DOCKET NUMBER: 377(913).6277P
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 215-875-8383
: TELEFAX: 215-875-8394
: INFORMATION FOR SEQ ID NO: 16:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 76 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: both
US-08-503-730-16
: Query Match 0.5%; Score 16; DB 1; Length 76;
: Best Local Similarity 87.5%; Pred. No. 2.2e+02;
: Matches 14; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 2165 gtcccaagccatcagc 2180
1:|||||||||:|||||
Db 5 GUCCCAAGCCCAUCCAGC 20

RESULT 42
US-08-507-634-13
: Sequence 13, Application US/08507634
: Patent No. 5849563
: GENERAL INFORMATION:
: APPLICANT: Miyata, Shohei
: APPLICANT: Ohshima, Atsushi
: APPLICANT: Inouye, Sumiko
: APPLICANT: Inouye, Masayori
: TITLE OF INVENTION: METHOD FOR SYNTHESIZING STABLE
: TITLE OF INVENTION: SINGLE-STRANDED CDNA IN EUKARYOTES BY MEANS OF A BACTERIAL
: NUMBER OF SEQUENCES: 24
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Weiser & Associates
: STREET: 230 South Fifteenth Street, Suite 500
: CITY: Philadelphia
: STATE: PA
: COUNTRY: USA
: ZIP: 19102
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ORIGINAL SOURCE:
INDIVIDUAL ISOLATE: INTRON 6 OF RAD50 GENOMIC SEQUENCE
S-08-687-080-70

Query Match 0.5%; Score 16; DB 2; Length 233;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
y 966 ttgtggtgtagaat 981
2 TTTGTGGTGTAGAT 17

RESULT 45
S-08-630-822A-97/c
Sequence 97, Application US/08630822A
Patent No. 5840695
GENERAL INFORMATION:
APPLICANT: FRANK, GLENN R.
APPLICANT: HUNTER, SHIRLEY WU
APPLICANT: WALLENFELS, LYNDIA
TITLE OF INVENTION: NOVEL ECTOPARASITE SALIVA PROTEINS
TITLE OF INVENTION: AND APPARATUS TO COLLECT SUCH PROTEINS
NUMBER OF SEQUENCES: 107
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sheridan Ross P.C.
STREET: 1700 Lincoln Street, Suite 3500
CITY: Denver
STATE: Colorado
COUNTRY: U.S.A.
ZIP: 80203

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/630,822A
FILING DATE: 11-APR-1996
CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:
NAME: CONNELL, GARY J.
REGISTRATION NUMBER: 32,020
REFERENCE/DOCKET NUMBER: 2618-17-C3
TELEPHONE: (303) 863-9700
TELEFAX: (303) 863-0223
INFORMATION FOR SEQ ID NO: 97:
SEQUENCE CHARACTERISTICS:
LENGTH: 252 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cdna

S-08-630-822A-97

Query Match 0.5%; Score 16; DB 2; Length 252;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

y 1478 agaagtcagtagccca 1493
195 AAGAGTCAGTACCCA 180

RESULT 46
S-09-005-069-97/c
Sequence 97, Application US/09005069
Patent No. 5932470
GENERAL INFORMATION:
APPLICANT: FRANK, GLENN R.

APPLICANT: HUNTER, SHIRLEY WU
APPLICANT: WALLENFELS, LYNDIA
TITLE OF INVENTION: NOVEL ECTOPARASITE SALIVA PROTEINS
TITLE OF INVENTION: AND APPARATUS TO COLLECT SUCH PROTEINS
NUMBER OF SEQUENCES: 107
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sheridan Ross P.C.
STREET: 1700 Lincoln Street, Suite 3500
CITY: Denver
STATE: Colorado
COUNTRY: U.S.A.
ZIP: 80203
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/005,069
FILING DATE:

CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/630,822
FILING DATE: 11-APR-1996
ATTORNEY/AGENT INFORMATION:
NAME: CONNELL, GARY J.
REGISTRATION NUMBER: 32,020
REFERENCE/DOCKET NUMBER: 2618-17-C3
TELEPHONE: (303) 863-9700
TELEFAX: (303) 863-0223
INFORMATION FOR SEQ ID NO: 97:
SEQUENCE CHARACTERISTICS:
LENGTH: 252 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cdna
US-09-005-069-97

Query Match 0.5%; Score 16; DB 2; Length 252;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1478 agaagtcagtagccca 1493
195 AAGAGTCAGTACCCA 180

RESULT 47
US-08-906-769-104/c
Sequence 104, Application US/08906769
Patent No. 6077687
GENERAL INFORMATION:
APPLICANT: Grieve, Robert B.
APPLICANT: Rushlow, Keith E.
APPLICANT: Wu Hunter, Shirley
APPLICANT: Frank, Glenn R.
APPLICANT: Stiegler, Gary
APPLICANT: Gaines, Patrick J.
APPLICANT: Silver, Gary
TITLE OF INVENTION: FLEA PROTEASE PROTEINS, NUCLEIC ACID
TITLE OF INVENTION: MOLECULES AND USES THEREOF
NUMBER OF SEQUENCES: 190
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sheridan Ross & McIntosh
STREET: 1700 Lincoln Street, Suite 3500
CITY: Denver
STATE: Colorado
COUNTRY: USA
ZIP: 80203
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/906.769
FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/639,075

FILING DATE: 24-APR-1996

ATTORNEY/AGENT INFORMATION:

NAME: Connell, Gary J.

REGISTRATION NUMBER: 32,020

REFERENCE/DOCKET NUMBER: 2618-25-C2

TELEPHONE: (303) 863-9700

TELEFAX: (303) 863-0223

INFORMATION FOR SEQ ID NO: 104:

SEQUENCE CHARACTERISTICS:

LENGTH: 252 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

FEATURE:

NAME/KEY: CDS

LOCATION: 1..251

OTHER INFORMATION: /note= "At pos. bp 4, change A to

OTHER INFORMATION: R. At pos. aa 2, substitute xaa."

US-08-906-769-104

Query Match 0.5%; Score 16; DB 3; Length 252;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1478 aagaagtcagtcacca 1493

Db 199 AGAAGTCAGTACCCA 184

RESULT 48

US-08-906-616-104/c

Sequence 104, Application US/08906616

Patent No. 6121035

GENERAL INFORMATION:

APPLICANT: Grieve, Robert B.

APPLICANT: Rushlow, Keith E.

APPLICANT: Wu Hunter, Shirley

APPLICANT: Frank, Glenn R.

APPLICANT: Stiegler, Gary

APPLICANT: Gaines, Patrick J.

APPLICANT: Silver, Gary

TITLE OF INVENTION: FLEA AMINOPEPTIDASE PROTEINS AND USES THEREOF

NUMBER OF SEQUENCES: 190

CORRESPONDENCE ADDRESS:

ADDRESSEE: Sheridan Ross P.C.

STREET: 1700 Lincoln Street, Suite 3500

CITY: Denver

STATE: Colorado

COUNTRY: USA

ZIP: 80203

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/906,616

FILING DATE: 05-AUG-1997

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: Connell, Gary J.
REGISTRATION NUMBER: 32,020
REFERENCE/DOCKET NUMBER: 2618-25-C2-3
TELEPHONE: (303) 863-9700
TELEFAX: (303) 863-0223
INFORMATION FOR SEQ ID NO: 104:
SEQUENCE CHARACTERISTICS:
LENGTH: 252 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..251
OTHER INFORMATION: /note= "At pos. bp 4, change A to
OTHER INFORMATION: R. At pos. aa 2, substitute xaa."
US-08-906-616-104

Query Match 0.5%; Score 16; DB 3; Length 252;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1478 aagaagtcagtcacca 1493

Db 199 AGAAGTCAGTACCCA 184

RESULT 49

US-08-817-795-104/c

Sequence 104, Application US/08817795

Patent No. 6139840

GENERAL INFORMATION:

APPLICANT: Grieve, Robert B.

APPLICANT: Rushlow, Keith E.

APPLICANT: Hunter, Shirley Wu

APPLICANT: Frank, Glenn R.

APPLICANT: Heath, Andrew W.

APPLICANT: Yamaka, Miles Yamanaka

APPLICANT: Arfsten, Ann

APPLICANT: Dale, Beverly

APPLICANT: Stiegler, Gary

TITLE OF INVENTION: USE OF PROTEASE INHIBITORS AND

TITLE OF INVENTION: PROTEASE VACCINES TO PROTECT ANIMALS FROM FLEA

TITLE OF INVENTION: INFESTATION, AND FLEA PROTEASE PROTEINS, NUCLEIC ACID

TITLE OF INVENTION: MOLECULES, AND USES THEREOF

NUMBER OF SEQUENCES: 119

CORRESPONDENCE ADDRESS:

ADDRESSEE: Sheridan Ross & McIntosh

STREET: 1700 Lincoln Street, Suite 3500

CITY: Denver

STATE: Colorado

COUNTRY: USA

ZIP: 80203

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/817,795

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: PCT/US95/14442

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Gary J. Connell

REGISTRATION NUMBER: 32,020

REFERENCE/DOCKET NUMBER:

TELECOMMUNICATION INFORMATION:

PCT-US95-14442A-104

Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1881 caggaagggtcgtgaga 1896
|||||
DB 184 CAGGAAGGCTGAGA 199

RESULT 57

PCT-US95-14792-7
Sequence 7, Application PC/TUS9514792
GENERAL INFORMATION:
APPLICANT: James Eberwine, Marc Dichter, Kevin Miyashiro
TITLE OF INVENTION: USE OF NEURITE LOCALIZED RNAs FOR
MEDICAL DIAGNOSIS AND THERAPEUTICS
NUMBER OF SEQUENCES: 28
CORRESPONDENCE ADDRESS:
ADDRESSEE: Jane Massey Licata, Esq.
STREET: 210 Lake Drive East, Suite 201
CITY: Cherry Hill
STATE: NJ USA
COUNTRY: USA
ZIP: 08002
COMPUTER READABLE FORM:
MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 Mb STORAGE
COMPUTER: IBM 486
OPERATING SYSTEM: WINDOWS FOR WORKGROUPS
SOFTWARE: WORDPERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14792
FILING DATE: Herewith
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Jane Massey Licata
REGISTRATION NUMBER: 32,257
REFERENCE/DOCKET NUMBER: PENN-0028
TELEPHONE: (609) 779-2400
TELEFAX: (609) 779-8488
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 489
TYPE: NUCLEIC ACID
STRANDEDNESS: SINGLE
TOPOLOGY: LINEAR
ANTI-SENSE: NO

PCT-US95-14792-7

Query Match 0.5%; Score 16; DB 4; Length 489;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1881 caggaagggtcgtgaga 1896
|||||
DB 184 CAGGAAGGCTGAGA 199

RESULT 58

PCT-US95-507-016-8
Sequence 8, Application US/08507016
Patent No. 5756460
GENERAL INFORMATION:
APPLICANT: EVANS, HELEN F.
APPLICANT: SHINE, JOHN
TITLE OF INVENTION: HUMAN GALANIN, CDNA CLONES ENCODING
HUMAN GALANIN AND A METHOD OF PRODUCING HUMAN GALANIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: ROTHWELL, FIGG, ERNST & KURZ

STREET: 555 THIRTEENTH STREET, N.W.
CITY: WASHINGTON
STATE: D. C.
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/507.016
FILING DATE: 25-JULY-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/108,733
FILING DATE: 03-SEP-1993
APPLICATION NUMBER: PCT/AU92/00097
FILING DATE: 06-MAR-1992
APPLICATION NUMBER: AU PK4953
FILING DATE: 06-MAR-1991

ATTORNEY/AGENT INFORMATION:
NAME: ERNST, BARBARA G.
REGISTRATION NUMBER: 30,377
REFERENCE/DOCKET NUMBER: 1871-117A
TELEPHONE: (202)783-6040
TELEFAX: (202)783-6031
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 575 base pairs
TYPE: nucleic acid
STRANDEDNESS: both
TOPOLOGY: linear
MOLECULE TYPE: DNA
HYPOTHETICAL: NO
FEATURE:
NAME/KEY: CDS
LOCATION: 14...385
US-08-507-016-8

Query Match 0.5%; Score 16; DB 1; Length 575;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2400 aagcgaggagctgcgc 2415
|||||
DB 200 AAGCGGAGCTGCGC 215

RESULT 59

PCT-US91-06418-4/c
Sequence 4, Application PC/TUS9106418
GENERAL INFORMATION:
APPLICANT: Oklahoma Medical Research, Foundation, et al
TITLE OF INVENTION: Antigens Associated with Polymyositis
and with Dermatomyositis
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: Kilpatrick & Cody
STREET: 100 Peachtree Street
CITY: Atlanta
STATE: Georgia
COUNTRY: US
ZIP: 30303
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US91/06418

FILING DATE: 19910905
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/579023
FILING DATE: 09-JUL-1990
ATTORNEY/AGENT INFORMATION:
NAME: Pabst, Patrea L.
REGISTRATION NUMBER: 31,284
REFERENCE/DOCKET NUMBER: OMR120
TELEPHONE: 404-572-6508
TELEFAX: 404-572-6555
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 578 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
FRAGMENT TYPE: N-terminal
ORIGINAL SOURCE:
ORGANISM: Homo sapien
STRAIN: JH2
TISSUE TYPE: Sera
CT-US91-06418-4

Query Match 0.5%; Score 16; DB 4; Length 578;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 955 ctggtgctgtttgt 970
|||||

22 CTGCTGCTGCTTTGT 7

RESULT 60

US-08-338-579A-94/c
Sequence 94, Application US/08338579A
Patent No. 6068975

GENERAL INFORMATION:

APPLICANT: Gilliam, T. Conrad
APPLICANT: Tanzi, Rudolph E.
TITLE OF INVENTION: ISOLATION AND USES OF A WILSON'S
DISEASE GENE
NUMBER OF SEQUENCES: 107
CORRESPONDENCE ADDRESS:

ADDRESSEE: Cooper & Dunham
STREET: 1185 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: United States of America
ZIP: 10036

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/338,579A
FILING DATE: June 17, 1996

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: White, John P.

REGISTRATION NUMBER: 28,678

REFERENCE/DOCKET NUMBER: 0575/44011-A-PCT-US

TELECOMMUNICATION INFORMATION:

TELEPHONE: (212) 278-0400

TELEFAX: (212) 391-0525

TELEX:

INFORMATION FOR SEQ ID NO: 94:

SEQUENCE CHARACTERISTICS:
LENGTH: 609 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 163..609
US-08-338-579A-94

Query Match 0.5%; Score 16; DB 3; Length 609;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2740 gggccaggaggtgcc 2755

|||||

Db 564 GGGCCAGGAGGCTGCC 549

RESULT 61

US-08-911-319A-2

Sequence 2, Application US/08911319A

Patent No. 5968798

GENERAL INFORMATION:

APPLICANT: Hillman, Jennifer L.

APPLICANT: Corley, Neil C.

APPLICANT: Shah, Purvi

TITLE OF INVENTION: HUMAN GLUTAREDOXIN BETA

NUMBER OF SEQUENCES: 6

CORRESPONDENCE ADDRESS:

ADDRESSEE: Incyte Pharmaceuticals, Inc.

STREET: 3174 Porter Dr.

CITY: Palo Alto

STATE: CA

COUNTRY: USA

ZIP: 94304

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/911,319A

FILING DATE: August 14, 1997

PRIOR APPLICATION DATA:

APPLICATION NUMBER:

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Muenzen, Colette C.

REGISTRATION NUMBER: 39,784

REFERENCE/DOCKET NUMBER: PF-0363 US

TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-855-0555

TELEFAX: 650-845-4166

INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:

LENGTH: 654 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

IMMEDIATE SOURCE:

LIBRARY: THPINOT03

CLONE: 2447829

US-08-911-319A-2

Query Match 0.5%; Score 16; DB 2; Length 654;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2402 gcgggagctgcgag 2417

25 101 GCGGAGCTGCGGAG 116
|||||

RESULT 62

Sequence 2, Application US/09352619
Patent No. 6084070
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Corley, Neil C.
APPLICANT: Shah, Purvi
TITLE OF INVENTION: HUMAN GLUTAREDOXIN BETA
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/352,619
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/911,319
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Muenzen, Colette C.
REGISTRATION NUMBER: 39,784
REFERENCE/DOCKET NUMBER: PF-0363 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 654 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: THPINOT03
CLONE: 2447829
S-09-352-619-2

Query Match 0.5%; Score 16; DB 3; Length 654;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

26 2402 GCGGAGCTGCGGAG 2417
|||||

26 101 GCGGAGCTGCGGAG 116
|||||

RESULT 63

Sequence 3, Application US/07789738
Patent No. 5824857
GENERAL INFORMATION:
APPLICANT: Beachy, Roger N.
APPLICANT: Bhattacharyya, Maitrayee
TITLE OF INVENTION: Plant Promoter
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Dennis R. Hoerner, Jr., Monsanto Co. BB4F
STREET: 700 Chesterfield Parkway No. 5824857th
CITY: St. Louis

STATE: Missouri
COUNTRY: USA
ZIP: 63198
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/789,738
FILING DATE: 19920330
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Hoerner Jr., Dennis R.
REGISTRATION NUMBER: 30,914
REFERENCE/DOCKET NUMBER: 38-21(10540)A
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314)537-6099
TELEFAX: (314)537-6047
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 714 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-07-789-738-3

Query Match 0.5%; Score 16; DB 1; Length 714;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1842 GTCCTGCACCATCA 1857
|||||

DB 484 GTCCTGCACCATCA 499
|||||

RESULT 64

US-07-789-738-5
Sequence 5, Application US/07789738
Patent No. 5824857
GENERAL INFORMATION:
APPLICANT: Beachy, Roger N.
APPLICANT: Bhattacharyya, Maitrayee
TITLE OF INVENTION: Plant Promoter
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Dennis R. Hoerner, Jr., Monsanto Co. BB4F
STREET: 700 Chesterfield Parkway No. 5824857th
CITY: St. Louis
STATE: Missouri
COUNTRY: USA
ZIP: 63198
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/789,738
FILING DATE: 19920330
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Hoerner Jr., Dennis R.
REGISTRATION NUMBER: 30,914
REFERENCE/DOCKET NUMBER: 38-21(10540)A
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314)537-6099
TELEFAX: (314)537-6047
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 773 base pairs

TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-07-789-738-5

Query Match 0.5%; Score 16; DB 1; Length 773;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1842 gtctgcacacacatca 1857
484 gtctgcacacacatca 499

RESULT 65

US-08-651-136C-15
Sequence 15, Application US/08651136C
Patent No. 6001639

GENERAL INFORMATION:

APPLICANT: Schuelein, Martin
APPLICANT: Andersen, Lene N.
APPLICANT: Lassen, Soren F.
APPLICANT: Kauppinen, Markus S.
APPLICANT: Lange, Lene
APPLICANT: Nielsen, Ruby I.
APPLICANT: Ihara, Michiko
TITLE OF INVENTION: Takagi, Shinobu
TITLE OF INVENTION: No. 6001639el Endoglucanases
NUMBER OF SEQUENCES: 109
CORRESPONDENCE ADDRESS:

ADDRESSEE: No. 6001639o No. 6001639disk of No. 6001639th America, Inc.
STREET: 405 Lexington Avenue, 64th Floor
CITY: New York
STATE: New York
COUNTRY: United States of America
ZIP: 10174-6401

COMPUTER READABLE FORM:

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/651,136C
FILING DATE: 21-MAY-1996

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:
NAME: Lambiris, Elias J.
REGISTRATION NUMBER: 33,728
REFERENCE/DOCKET NUMBER: 4366.200-US
TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-867-0123

TELEFAX: 212-878-9655

INFORMATION FOR SEQ ID NO: 15:

SEQUENCE CHARACTERISTICS:

LENGTH: 808 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

FEATURE:

NAME/KEY: CDS

LOCATION: 37..714

US-08-651-136C-15

Query Match 0.5%; Score 16; DB 3; Length 808;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1409 tcccaacttcacagcag 1424
|||||

Db 577 TCCCACTTCACAG 592

RESULT 66

US-08-906-769-128/c
Sequence 128, Application US/08906769
Patent No. 6077687

GENERAL INFORMATION:

APPLICANT: Grieve, Robert B.
APPLICANT: Rushlow, Keith E.
APPLICANT: Wu Hunter, Shirley
APPLICANT: Frank, Glenn R.
APPLICANT: Stiegler, Gary
APPLICANT: Gaines, Patrick J.
APPLICANT: Silver, Gary
TITLE OF INVENTION: FLEA PROTEASE PROTEINS, NUCLEIC ACID
TITLE OF INVENTION: MOLECULES AND USES THEREOF
NUMBER OF SEQUENCES: 190
CORRESPONDENCE ADDRESS:

ADDRESSEE: Sheridan Ross & McIntosh
STREET: 1700 Lincoln Street, Suite 3500
CITY: Denver
STATE: Colorado
COUNTRY: USA
ZIP: 80203

COMPUTER READABLE FORM:

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/906,769
FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/639,075
FILING DATE: 24-APR-1996
ATTORNEY/AGENT INFORMATION:

NAME: Connell, Gary J.

REGISTRATION NUMBER: 32,020

REFERENCE/DOCKET NUMBER: 2618-25-C2

TELECOMMUNICATION INFORMATION:

TELEPHONE: (303) 863-9700

TELEFAX: (303) 863-0223

INFORMATION FOR SEQ ID NO: 128:

SEQUENCE CHARACTERISTICS:

LENGTH: 815 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

FEATURE:

NAME/KEY: CDS

LOCATION: 1..762

OTHER INFORMATION:

OTHER INFORMATION: M; at 454, change G to V; at 456, G to V; at 457, A to M;

OTHER INFORMATION: 460, A to R; at 470, G to S; at 493, A to R. At pos. aa 1

OTHER INFORMATION: 136, 152, 153, 154, 157 and 165, substitute xaa.

US-08-906-769-128

Query Match 0.5%; Score 16; DB 3; Length 815;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1478 aagaagtcagtcacca 1493
|||||

Db 181 AAGAGTCAGTACCCA 166

RESULT 67

US-08-906-616-128/c
Sequence 128, Application US/08906616

Patent No. 6121035
GENERAL INFORMATION:
APPLICANT: Grieve, Robert B.
APPLICANT: Rushlow, Keith E.
APPLICANT: Wu Hunter, Shirley
APPLICANT: Frank, Glenn R.
APPLICANT: Stiegler, Gary
APPLICANT: Gaines, Patrick J.
APPLICANT: Silver, Gary
TITLE OF INVENTION: FLEA AMINOPEPTIDASE PROTEINS AND USES THEREOF
NUMBER OF SEQUENCES: 190
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sheridan Ross P.C.
STREET: 1700 Lincoln Street, Suite 3500
CITY: Denver
STATE: Colorado
COUNTRY: USA
ZIP: 80203
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/906.616
FILING DATE: 05-AUG-1997
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Connell, Gary J.
REGISTRATION NUMBER: 32,020
REFERENCE/DOCKET NUMBER: 2618-25-C2-3
TELEPHONE: (303) 863-9700
TELEFAX: (303) 863-0223
INFORMATION FOR SEQ ID NO: 128:
SEQUENCE CHARACTERISTICS:
LENGTH: 815 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..762
OTHER INFORMATION: /note= "At pos. bp 453, change A to M; at 454, change G to V; at 456, G to V; at 457, A to M; at 460, A to R; at 470, G to S; at 493, A to R. At pos. aa 120, 136, 152, 153, 154, 157 and 165, substitute Xaa."
US-08-906-616-128

Query Match 0.5%; Score 16; DB 3; Length 815;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1478 aagaagtcagtagcca 1493
Db 181 AAGAAGTCAGTACCCA 166
|||||
RESULT 68
US-08-639-075A-128/c
Sequence 128, Application US/08639075A
Patent No. 6150125
GENERAL INFORMATION:
APPLICANT: Grieve, Robert B.
APPLICANT: Rushlow, Keith E.
APPLICANT: Wu Hunter, Shirley
APPLICANT: Frank, Glenn R.
APPLICANT: Stiegler, Gary
APPLICANT: Gaines, Patrick J.
APPLICANT: Silver, Gary
TITLE OF INVENTION: FLEA PROTEASE, PROTEINS, NUCLEIC ACID

TITLE OF INVENTION: MOLECULES AND USES THEREOF
NUMBER OF SEQUENCES: 190
CORRESPONDENCE ADDRESS:
ADDRESSEE: Sheridan Ross & McIntosh
STREET: 1700 Lincoln Street, Suite 3500
CITY: Denver
STATE: Colorado
COUNTRY: USA
ZIP: 80203
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/639.075A
FILING DATE: 24-APR-1996
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Connell, Gary J.
REGISTRATION NUMBER: 32,020
REFERENCE/DOCKET NUMBER: 2618-25-C2
TELEPHONE: (303) 863-9700
TELEFAX: (303) 863-0223
INFORMATION FOR SEQ ID NO: 128:
SEQUENCE CHARACTERISTICS:
LENGTH: 815 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..762
OTHER INFORMATION: /note= "At pos. bp 453, change A to M; at 454, change G to V; at 456, G to V; at 457, A to M; at 460, A to R; at 470, G to S; at 493, A to R. At pos. aa 136, 152, 153, 154, 157 and 165, substitute Xaa."
US-08-639-075A-128

Query Match 0.5%; Score 16; DB 3; Length 815;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1478 aagaagtcagtagcca 1493
Db 181 AAGAAGTCAGTACCCA 166
|||||
RESULT 69
US-08-651-136C-19
Sequence 19, Application US/08651136C
Patent No. 6001639
GENERAL INFORMATION:
APPLICANT: Schulein, Martin
APPLICANT: Andersen, Lene N.
APPLICANT: Lassen, Soren F.
APPLICANT: Kauppinen, Markus S.
APPLICANT: Lange, Lene
APPLICANT: Nielsen, Ruby I.
APPLICANT: Ihara, Michiko
APPLICANT: Takagi, Shinobu
TITLE OF INVENTION: No. 6001639e1 Endoglucanases
NUMBER OF SEQUENCES: 109
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 60016390 No. 6001639disk of No. 6001639th America, Inc.
STREET: 405 Lexington Avenue, 64th Floor
CITY: New York
STATE: New York
COUNTRY: United States of America
ZIP: 10174-6401

```
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/651,136C
FILING DATE: 21-MAY-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Lambiris, Elias J.
REGISTRATION NUMBER: 33,728
REFERENCE/DOCKET NUMBER: 4366.200-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-867-0123
TELEFAX: 212-878-9655
INFORMATION FOR SEQ ID NO: 19:
SEQUENCE CHARACTERISTICS:
LENGTH: 1031 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDS
FEATURE:
NAME/KEY: CDS
LOCATION: 11..889
US-08-651-136C-19

Query Match 0.5%; Score 16; DB 3; Length 1031;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1409 tcccaacttcagcag 1424
|||||
536 TCCCAACTTCCAGCAG 551

RESULT 70
US-08-651-136C-17
Sequence 17, Application US/08651136C
Patent No. 6001639
GENERAL INFORMATION:
APPLICANT: Schulein, Martin
APPLICANT: Andersen, Lene N.
APPLICANT: Lassen, Soren F.
APPLICANT: Kauppinen, Markus S.
APPLICANT: Lange, Lene
APPLICANT: Nielsen, Ruby I.
APPLICANT: Ihara, Michiko
APPLICANT: Takagi, Shinobu
TITLE OF INVENTION: No. 6001639el Endoglucanases
NUMBER OF SEQUENCES: 109
CORRESPONDENCE ADDRESS:
ADDRESS: No. 6001639o No. 6001639disk of No. 6001639th America, Inc.
STREET: 405 Lexington Avenue, 64th Floor
CITY: New York
STATE: New York
COUNTRY: United States of America
ZIP: 10174-6401
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/651,136C
FILING DATE: 21-MAY-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Lambiris, Elias J.
REGISTRATION NUMBER: 33,728
REFERENCE/DOCKET NUMBER: 4366.200-US

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/651,136C
FILING DATE: 21-MAY-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Lambiris, Elias J.
REGISTRATION NUMBER: 33,728
REFERENCE/DOCKET NUMBER: 4366.200-US

TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-867-0123
TELEFAX: 212-878-9655
INFORMATION FOR SEQ ID NO: 17:
SEQUENCE CHARACTERISTICS:
LENGTH: 1048 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 13..906
US-08-651-136C-17

Query Match 0.5%; Score 16; DB 3; Length 1048;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1409 tcccaacttcagcag 1424
|||||
553 TCCCAACTTCCAGCAG 568

RESULT 71
US-08-933-750C-81/c
Sequence 81, Application US/08933750C
Patent No. 5932442
GENERAL INFORMATION:
APPLICANT: Lal, Preeti
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
APPLICANT: Shah, Purvi
APPLICANT: Au-Young, Janice
APPLICANT: Yue, Henry
APPLICANT: Guegler, Karl J.
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: HUMAN REGULATORY MOLECULES
NUMBER OF SEQUENCES: 98
CORRESPONDENCE ADDRESS:
ADDRESS: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/933,750C
FILING DATE: September 23, 1997
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0356 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 81:
SEQUENCE CHARACTERISTICS:
LENGTH: 1152 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
```

IMMEDIATE SOURCE:
LIBRARY: BLADNOT03
CLONE: 1602473
US-08-933-750C-81

Query Match 0.5%; Score 16; DB 2; Length 1152;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 2168 ccaagccatcagcggtg 2183
DB 928 CCAAGCCATCAGCGGTG 913
|||||

RESULT 72

US-09-234-613-81/c
Sequence 81, Application US/09234613
Patent No. 6132973
GENERAL INFORMATION:
APPLICANT: Lal, Preeti
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
APPLICANT: Shah, Purvi
APPLICANT: Au-Young, Janice
APPLICANT: Yue, Henry
APPLICANT: Guegler, Karl J.
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: HUMAN REGULATORY MOLECULES
NUMBER OF SEQUENCES: 98
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/234,613
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/933,750
FILING DATE: September 23, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0356 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:

INFORMATION FOR SEQ ID NO: 81:
SEQUENCE CHARACTERISTICS:
LENGTH: 1152 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: BLADNOT03
CLONE: 1602473
US-09-234-613-81

Query Match 0.5%; Score 16; DB 3; Length 1152;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2168 ccaagccatcagcggtg 2183
DB 928 CCAAGCCATCAGCGGTG 913
|||||

RESULT 73

US-08-739-485-4
Sequence 4, Application US/08739485
Patent No. 5863898
GENERAL INFORMATION:
APPLICANT: Goli, Surya K.
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
TITLE OF INVENTION: NOVEL HUMAN LIM PROTEINS
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: US
ZIP: 94304

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/739,485
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0142 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:

INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1225 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: Consensus
CLONE: Consensus
US-08-739-485-4

Query Match 0.5%; Score 16; DB 2; Length 1225;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1835 ccaggagggtcctgcac 1850
DB 441 CCAGGAGGTCTCTGCAC 456
|||||

RESULT 74

US-08-360-758-1/c
Sequence 1, Application US/08360758
Patent No. 6074863
GENERAL INFORMATION:
APPLICANT: Svendsen, Allan
APPLICANT: Pathar, Shankant A
APPLICANT: Egel-Mitani, Michi
APPLICANT: Borch, Kim
APPLICANT: Clausen, Ib G

APPLICANT: Hansen, Mogens T
TITLE OF INVENTION: C. ANTARCTICA LIPASE AND LIPASE VARIANTS
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 60748630 No. 6074863disk of No. 6074863th America, Inc.
STREET: 405 Lexington Avenue, 64th Floor
CITY: New York
STATE: New York
COUNTRY: United States of America
ZIP: 10174-6401
COMPUTER READABLE FORM:
MEDIUM TYPE: Tape
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/360,758
FILING DATE: 22-DEC-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DK PCT/DK93/00225
FILING DATE: 03-JUN-1993
ATTORNEY/AGENT INFORMATION:
NAME: Lambiris, Elias J.
REGISTRATION NUMBER: 33,728
REFERENCE/DOCKET NUMBER: 3748.204-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-867-0123
TELEFAX: 212-878-9655
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1329 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-360-758-1

Query Match 0.5%; Score 16; DB 3; Length 1329;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1143 accagcactgtgcc 1158
|||||
DB 1245 ACCGACCTGTGCTCC 1230

RESULT 75
US-08-889-425-3/C
Sequence 3, Application US/08889425
Patent No. 6153403
GENERAL INFORMATION:
APPLICANT: Lim, Bing
APPLICANT: Adra, Chaker N.
TITLE OF INVENTION: A Lysosomal-Associated Multispanning
TITLE OF INVENTION: Membrane Protein, LAPTM5 and a Nucleic Acid Encoding
TITLE OF INVENTION: LAPTM5
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
STREET: Two Militia Drive
CITY: Lexington
STATE: Massachusetts
COUNTRY: USA
ZIP: 02173
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/889,425

APPLICANT: Hansen, Mogens T
TITLE OF INVENTION: C. ANTARCTICA LIPASE AND LIPASE VARIANTS
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: No. 60748630 No. 6074863disk of No. 6074863th America, Inc.
STREET: 405 Lexington Avenue, 64th Floor
CITY: New York
STATE: New York
COUNTRY: United States of America
ZIP: 10174-6401
COMPUTER READABLE FORM:
MEDIUM TYPE: Tape
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/360,758
FILING DATE: 22-DEC-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DK PCT/DK93/00225
FILING DATE: 03-JUN-1993
ATTORNEY/AGENT INFORMATION:
NAME: Lambiris, Elias J.
REGISTRATION NUMBER: 33,728
REFERENCE/DOCKET NUMBER: 3748.204-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-867-0123
TELEFAX: 212-878-9655
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1329 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-360-758-1

Query Match 0.5%; Score 16; DB 3; Length 1329;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1143 accagcactgtgcc 1158
|||||
DB 1245 ACCGACCTGTGCTCC 1230

RESULT 75
US-08-889-425-3/C
Sequence 3, Application US/08889425
Patent No. 6153403
GENERAL INFORMATION:
APPLICANT: Lim, Bing
APPLICANT: Adra, Chaker N.
TITLE OF INVENTION: A Lysosomal-Associated Multispanning
TITLE OF INVENTION: Membrane Protein, LAPTM5 and a Nucleic Acid Encoding
TITLE OF INVENTION: LAPTM5
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
STREET: Two Militia Drive
CITY: Lexington
STATE: Massachusetts
COUNTRY: USA
ZIP: 02173
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/889,425

FILING DATE: 08-JUL-1997
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: Brook, David E.
REGISTRATION NUMBER: 22,592
REFERENCE/DOCKET NUMBER: BIH96-09pa
TELECOMMUNICATION INFORMATION:
TELEPHONE: (781) 861-6240
TELEFAX: (781) 861-9540
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1333 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 58..840
US-08-889-425-3

Query Match 0.5%; Score 16; DB 3; Length 1333;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 534 gaactggctgtgcggc 549
|||||
DB 606 GAACTGGCTGTGCGGC 591

RESULT 76
US-09-032-372-9/C
Sequence 9, Application US/09032372
Patent No. 6008337
GENERAL INFORMATION:
APPLICANT: Bandman, Olga
APPLICANT: Hillman, Jennifer L.
APPLICANT: Corley, Neil C.
APPLICANT: Guegler, Karl J.
APPLICANT: Yue, Henry
APPLICANT: Lal, Preeti
TITLE OF INVENTION: CELL CYCLE RELATED PROTEINS
NUMBER OF SEQUENCES: 13
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/032,372
FILING DATE: Herewith
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0478-US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:

LENGTH: 1341 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: BRAITUT21
CLONE: 2522306
3-09-032-372-9

Query Match 0.5%; Score 16; DB 3; Length 1341;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2624 cacggtcccccaggag 2639

DB 794 CACGGTCCCCCAGGAG 779

RESULT 77

US-08-374-155A-7
Sequence 7, Application US/08374155A
Patent No. 5786140

GENERAL INFORMATION:

APPLICANT: Mattes, Ralf
APPLICANT: Klein, Kathrin
APPLICANT: Schiweck, Hubert
APPLICANT: Kunz, Markwart
APPLICANT: Munir, Mohammed
TITLE OF INVENTION: Preparation of Acariogenic Sugar
TITLE OF INVENTION: Substitutes
NUMBER OF SEQUENCES: 26
CORRESPONDENCE ADDRESS:

ADDRESSEE: Finnegan, Henderson, Farabow, Garrett &

ADDRESSEE: Dunner

STREET: 1300 I Street, N.W.

CITY: Washington

STATE: D.C.

COUNTRY: USA

ZIP: 20005-3315

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC Compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/374,155A

FILING DATE: 18-JAN-1995

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Forman, David S

REGISTRATION NUMBER: 33,694

REFERENCE/DOCKET NUMBER: 05638.0006-00000

TELECOMMUNICATION INFORMATION:

TELEPHONE: (202) 408-4000

TELEFAX: (202) 408-4400

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 1362 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-08-374-155A-7

Query Match 0.5%; Score 16; DB 1; Length 1362;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2213 gctgaaccacttcagc 2228

DB 594 GCTGAACCACCTTCAGC 609

RESULT 78

US-08-785-396-7

Sequence 7, Application US/08785396

Patent No. 5985622

GENERAL INFORMATION:

APPLICANT: Mattes, Ralf

APPLICANT: Klein, Kathrin

APPLICANT: Schiweck, Hubert

APPLICANT: Kunz, Markwart

APPLICANT: Munir, Mohammed

TITLE OF INVENTION: Preparation of Acariogenic Sugar

TITLE OF INVENTION: Substitutes

NUMBER OF SEQUENCES: 26

CORRESPONDENCE ADDRESS:

ADDRESSEE: Finnegan, Henderson, Farabow, Garrett &

ADDRESSEE: Dunner

STREET: 1300 I Street, N.W.

CITY: Washington

STATE: D.C.

COUNTRY: USA

ZIP: 20005-3315

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC Compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/785,396

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/374,155

FILING DATE: 18-JAN-1995

ATTORNEY/AGENT INFORMATION:

NAME: Forman, David S

REGISTRATION NUMBER: 33,694

REFERENCE/DOCKET NUMBER: 05638.0006-00000

TELECOMMUNICATION INFORMATION:

TELEPHONE: (202) 408-4000

TELEFAX: (202) 408-4400

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 1362 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-08-785-396-7

Query Match 0.5%; Score 16; DB 2; Length 1362;

Best Local Similarity 100.0%; Pred. No. 2.2e+02;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2213 gctgaaccacttcagc 2228

DB 594 GCTGAACCACCTTCAGC 609

RESULT 79

US-08-458-023B-1/C

Sequence 1, Application US/08458023B

Patent No. 5667990

GENERAL INFORMATION:

APPLICANT: Berka, Randy M.

APPLICANT: Yoder, Wendy

APPLICANT: Takagi, Shinobu

APPLICANT: Boomnathan, Karuppan C.

TITLE OF INVENTION: ASPERGILLUS EXPRESSION SYSTEM

NUMBER OF SEQUENCES: 16

CORRESPONDENCE ADDRESS:

ADDRESSEE: No. 5667990o No. 5667990disk of No. 5667990th America, Inc.

STREET: 405 Lexington Avenue

CITY: New York

STATE: New York

COUNTRY: USA

ZIP: 10174-6201

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/458,023B

FILING DATE: 01-JUN-1995

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Lowrey Dr., Karen A.

REGISTRATION NUMBER: 31,274

REFERENCE/DOCKET NUMBER: 4086.010-US

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-867-0123

TELEFAX: 212-878-9655

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1389 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

HYPOTHETICAL: NO

ANTI-SENSE: NO

ORIGINAL SOURCE:

ORGANISM: Candida antarctica

INDIVIDUAL ISOLATE: DSM 3855

FEATURE:

NAME/KEY: CDS

LOCATION: 1..1389

5-08-458-023B-1

Query Match

Best Local Similarity 0.5%; Score 16; DB 1; Length 1389;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1143 acccagcacttggtcc 1158

|||||

DB 1305 ACCCAGCACTTGCTCC 1290

RESULT 80

US-09-111-556A-1/c

Sequence 1, Application US/09111556A

Patent No. 6020180

GENERAL INFORMATION:

APPLICANT: Svendsen, Allan

APPLICANT: Pathar, Shamkant A

APPLICANT: Egel-Mitani, Michi

APPLICANT: Borch, Kim

APPLICANT: Clausen, Ib G

APPLICANT: Hansen, Mogens T

TITLE OF INVENTION: C. ANTARCTICA LIPASE AND LIPASE VARIANTS

NUMBER OF SEQUENCES: 11

CORRESPONDENCE ADDRESS:

ADDRESSEE: No. 6020180o No. 6020180disk of No. 6020180th America, Inc.

STREET: 405 Lexington Avenue, 64th Floor

CITY: New York

STATE: New York

COUNTRY: United States of America

ZIP: 10174-6401

COMPUTER READABLE FORM:

MEDIUM TYPE: Tape

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/111,556A

FILING DATE: 22-DEC-1994

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: DK PCT/DK93/00225

FILING DATE: 03-JUN-1993

ATTORNEY/AGENT INFORMATION:

NAME: Lambiris, Elias J.

REGISTRATION NUMBER: 33,728

REFERENCE/DOCKET NUMBER: 3748.214-US

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-867-0123

TELEFAX: 212-878-9655

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1389 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-09-111-556A-1

Query Match

Best Local Similarity 0.5%; Score 16; DB 3; Length 1389;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1143 acccagcacttggtcc 1158

|||||

DB 1305 ACCCAGCACTTGCTCC 1290

RESULT 81

US-08-135-510-4/c

Sequence 4, Application US/08135510

Patent No. 5420028

GENERAL INFORMATION:

APPLICANT: CHIANG, John Young Ling

TITLE OF INVENTION: Truncated Human Cholesterol

7a-Hydroxylase, Method of Production and Use Thereof

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Foley & Lardner

STREET: 3000 K Street, N.W., Suite 500

CITY: Washington, D.C.

COUNTRY: USA

ZIP: 20007-5109

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/135,510

FILING DATE: 13-OCT-1993

CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:

NAME: SANDERCOCK, Colin G.

REGISTRATION NUMBER: 31,298

REFERENCE/DOCKET NUMBER: 18748/176 HOCE

TELECOMMUNICATION INFORMATION:

TELEPHONE: (202)672-5300

TELEFAX: (202)672-5399

TELEX: 904136

INFORMATION FOR SEQ ID NO: 4:

SEQUENCE CHARACTERISTICS:

LENGTH: 1524 base pairs

TYPE: nucleic acid

STRANDEDNESS: double

TOPOLOGY: linear

US-08-135-510-4

NUMBER OF SEQUENCES: 16
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: CHRISTINE E. CARTY - MERCK & CO., INC.
 STREET: 126 EAST LINCOLN AVENUE - P.O. BOX 2000
 CITY: RAHWAY
 STATE: NJ
 COUNTRY: US
 ZIP: 07065-0907
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: DOS
 SOFTWARE: FastSEQ Version 1.5
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/409,122
 FILING DATE:
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/408,669
 FILING DATE: 22-MAR-1995
 ATTORNEY/AGENT INFORMATION:
 NAME: CARTY, CHRISTINE E
 REGISTRATION NUMBER: 36,099
 REFERENCE/DOCKET NUMBER: 19425
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 908-594-6734
 TELEFAX: 908-594-4720
 TELEX:
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1524 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: CDNA
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 FRAGMENT TYPE:
 ORIGINAL SOURCE:
 US-08-409-122-1

Query Match 0.5%; Score 16; DB 1; Length 1524;
 Best Local Similarity 100.0%; Pred. No. 2.2e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 227 cctgcagggtggtggca 242
 |||||
 Db 157 CCTGCAGGTGTTGGCA 172
 RESULT 85
 US-08-408-669-1
 Sequence 1, Application US/08408669
 Patent No. 5840306
 GENERAL INFORMATION:
 APPLICANT: HOFMANN, KATHRYN J.
 APPLICANT: JANSEN, KATHRYN U.
 APPLICANT: NEEPER, MICHAEL P.
 TITLE OF INVENTION: DNA ENCODING HUMAN PAPILLOMAVIRUS TYPE 18
 NUMBER OF SEQUENCES: 16
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: CHRISTINE E. CARTY - MERCK & CO., INC.
 STREET: 126 EAST LINCOLN AVENUE - P.O. BOX 2000
 CITY: RAHWAY
 STATE: NJ
 COUNTRY: US
 ZIP: 07065-0907
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: DOS
 SOFTWARE: FastSEQ Version 1.5

CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/408,669
 FILING DATE: 22-MAR-1995
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER:
 FILING DATE:
 ATTORNEY/AGENT INFORMATION:
 NAME: CARTY, CHRISTINE E
 REGISTRATION NUMBER: 36,099
 REFERENCE/DOCKET NUMBER: 19424
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 908-594-6734
 TELEFAX: 908-594-4720
 TELEX:
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1524 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: CDNA
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 FRAGMENT TYPE:
 ORIGINAL SOURCE:
 US-08-408-669-1

Query Match 0.5%; Score 16; DB 2; Length 1524;
 Best Local Similarity 100.0%; Pred. No. 2.2e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 227 cctgcagggtggtggca 242
 |||||
 Db 157 CCTGCAGGTGTTGGCA 172

RESULT 86
 US-08-477-952-4/c
 Sequence 4, Application US/08477952
 Patent No. 5851780
 GENERAL INFORMATION:
 APPLICANT: CHIANG, John Young Ling
 TITLE OF INVENTION: Genomic DNA of Human Cholesterol
 NUMBER OF SEQUENCES: 11
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Foley & Lardner
 STREET: 3000 K Street, N.W., Suite 500
 CITY: Washington
 STATE: D.C.
 COUNTRY: USA
 ZIP: 20007-5109
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC Compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent in Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/477,952
 FILING DATE: 07-JUN-1995
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/361,458
 FILING DATE: 21-DEC-1994
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/135,511
 FILING DATE: 13-OCT-1993
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/135,488
 FILING DATE: 13-OCT-1993
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/135,510

FILING DATE: 13-OCT-1993
ATTORNEY/AGENT INFORMATION:
NAME: SANDERCOCK, COLIN G.
REGISTRATION NUMBER: 31,298
REFERENCE/DOCKET NUMBER: 18748/221 HOCE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1524 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
US-08-477-952-4

Query Match 0.5%; Score 16; DB 2; Length 1524;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 499 caatcaaaatatttc 514
|||||
Db 252 CAATCAAAATATTTC 237

RESULT 87
US-08-386-727-5/c
Sequence 5, Application US/08386727
Patent No. 5792647
GENERAL INFORMATION:
APPLICANT: ROSEMAN, SAUL
APPLICANT: BASSLER, BONNIE
APPLICANT: KEYHANI, NEMAT O.
APPLICANT: CHITLARU, EDITH
APPLICANT: ROWE, CHRIS
APPLICANT: YU, CHARLES
TITLE OF INVENTION: BACTERIAL CATABOLISM OF CHITIN
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: CUSHMAN, DARBY & CUSHMAN
STREET: 1100 NEW YORK AVENUE, N.W.
CITY: WASHINGTON
STATE: D.C.
COUNTRY: USA
ZIP: 20005
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/386.727
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: HOBBS, ANN S.
REGISTRATION NUMBER: 36,830
REFERENCE/DOCKET NUMBER: 4130/206916
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-861-3000
TELEFAX: 202-822-0944
TELEX: 6714627 CUSH
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 1713 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-386-727-5

FILING DATE: 13-FEB-1996
ATTORNEY/AGENT INFORMATION:
NAME: Haille, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07662/005001
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 678-5070
TELEFAX: (619) 678-5099
TELEX:
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 1713 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-600-452A-5

Query Match 0.5%; Score 16; DB 2; Length 1713;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1176 tcagttcacaccccttc 1191
|||||
Db 241 TCAGTTCACACCTTC 226

RESULT 88
US-08-600-452A-5/c
Sequence 5, Application US/08600452A
Patent No. 5985644
GENERAL INFORMATION:
APPLICANT: ROSEMAN, SAUL
APPLICANT: BASSLER, BONNIE
APPLICANT: KEYHANI, NEMAT O.
APPLICANT: CHITLARU, EDITH
APPLICANT: ROWE, CHRIS
APPLICANT: YU, CHARLES
TITLE OF INVENTION: BACTERIAL CATABOLISM OF CHITIN
NUMBER OF SEQUENCES: 8
CORRESPONDENCE ADDRESS:
ADDRESSEE: FISH & RICHARDSON P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/600.452A
FILING DATE: 13-FEB-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Haille, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07662/005001
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 678-5070
TELEFAX: (619) 678-5099
TELEX:
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 1713 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-600-452A-5

Query Match 0.5%; Score 16; DB 2; Length 1713;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1176 tcagttcacaccccttc 1191
|||||
Db 241 TCAGTTCACACCTTC 226

RESULT 89
US-08-467-948A-3/c
Sequence 1, Application US/08467948A
Patent No. 5998164
GENERAL INFORMATION:
APPLICANT: LI, YI
APPLICANT: CAO, LIANG

APPLICANT: NI, JIAN
APPLICANT: GENTZ, REINER
APPLICANT: BULT, CAROL J.
APPLICANT: SUTTON III, GRANGER G.
APPLICANT: ROSEN, CRAIG A.
TITLE OF INVENTION: Polynucleotides Encoding Human G-Protein
TITLE OF INVENTION: Coupled Receptor GPR2
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 NEW YORK AVE., NW, SUITE 600
CITY: WASHINGTON
STATE: DC
COUNTRY: USA
ZIP: 20005

COMPUTER READABLE FORM:
MEDIUM TYPE: FLOPPY DISK
COMPUTER: IBM PC COMPATIBLE
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PATENTIN RELEASE #1.0, VERSION #1.30
CURRENT APPLICATION NUMBER: US/08/467,948A

FILING DATE: 06-JUN-1995
CLASSIFICATION: 435

PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/04079

FILING DATE: 30-MAR-1995
ATTORNEY/AGENT INFORMATION:
NAME: STEFFE, ERIC K.

REGISTRATION NUMBER: 36,688
REFERENCE/DOCKET NUMBER: 1488.1140003/EKS/KLM
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1713 base pairs

TYPE: nucleic acid

STRANDEDNESS: both

TOPOLOGY: both

MOLECULE TYPE: cdna

FEATURE:

NAME/KEY: CDS

LOCATION: 116..1003

US-08-467-948A-1

Query Match 0.5%; Score 16; DB 2; Length 1713;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 824 agcaaaaggagatgggc 839
|||||
D 409 AGCAAGGAGATGGGC 394

RESULT 90

US-08-467-947A-1/c
Sequence 1, Application US/08467947A
Patent No. 6090575

GENERAL INFORMATION:

APPLICANT: LI, YI

APPLICANT: CAO, LIANG

APPLICANT: NI, JIAN

APPLICANT: GENTZ, REINER

APPLICANT: BULT, CAROL J.

APPLICANT: SUTTON III, GRANGER G.

APPLICANT: ROSEN, CRAIG A.

TITLE OF INVENTION: Polynucleotides Encoding Human G-Protein

TITLE OF INVENTION: Coupled Receptor GPR1

NUMBER OF SEQUENCES: 30

CORRESPONDENCE ADDRESS:

ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.

STREET: 1100 NEW YORK AVE., NW, SUITE 600
CITY: WASHINGTON
STATE: DC
COUNTRY: USA
ZIP: 20005
COMPUTER READABLE FORM:
MEDIUM TYPE: FLOPPY DISK
COMPUTER: IBM PC COMPATIBLE
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PATENTIN RELEASE #1.0, VERSION #1.30
CURRENT APPLICATION NUMBER: US/08/467,947A

FILING DATE: 06-JUN-1995

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: PCT/US95/04079

FILING DATE: 30-MAR-1995

ATTORNEY/AGENT INFORMATION:

NAME: STEFFE, ERIC K.

REGISTRATION NUMBER: 36,688

REFERENCE/DOCKET NUMBER: 1488.1140002/EKS/KLM

TELEPHONE: 202-371-2600

TELEFAX: 202-371-2540

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1713 base pairs

TYPE: nucleic acid

STRANDEDNESS: both

TOPOLOGY: both

MOLECULE TYPE: cdna

FEATURE:

NAME/KEY: CDS

LOCATION: 116..1003

US-08-467-947A-1

Query Match 0.5%; Score 16; DB 3; Length 1713;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 824 agcaaaaggagatgggc 839
|||||

DB 409 AGCAAGGAGATGGGC 394

RESULT 91

US-07-688-352C-13

Sequence 13, Application US/07688352C

Patent No. 5527896

GENERAL INFORMATION:

APPLICANT: Wigler, Michael H.

APPLICANT: Colicelli, John J.

TITLE OF INVENTION: Cloning by Complementation and Related

TITLE OF INVENTION: Processes

NUMBER OF SEQUENCES: 57

CORRESPONDENCE ADDRESS:

ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &

ADDRESSEE: Bicknell

STREET: Two First National Plaza, 20 South Clark

STREET: Street

CITY: Chicago

STATE: Illinois

COUNTRY: USA

ZIP: 60603

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/07/688,352C

FILING DATE: 19910419

```

: CLASSIFICATION: 435
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/511,715
: FILING DATE: 20-APR-1990
: ATTORNEY/AGENT INFORMATION:
: NAME: Borun, Michael F.
: REGISTRATION NUMBER: 25447
: REFERENCE/DOCKET NUMBER: 27805/30197
: TELEPHONE: (312) 346-5750
: TELEFAX: (312) 984-9740
: TELEX: 25-3856
: INFORMATION FOR SEQ ID NO: 13:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1721 base pairs
: TYPE: NUCLEIC ACID
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: CDNA
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 60..1274
: s-07-688-352C-13

Query Match 0.5%; Score 16; DB 1; Length 1721;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2731 cacagccgcgggccag 2746
DB 266 CACAGCCGCGGCCAG 281

RESULT 92
: s-08-474-379C-13
: Sequence 13, Application US/08474379C
: Patent No. 5977305
: GENERAL INFORMATION:
: APPLICANT: Wigler, Michael H.
: APPLICANT: Colicelli, John J.
: TITLE OF INVENTION: CLONING BY COMPLEMENTATION AND RELATED
: TITLE OF INVENTION: PROCESSES
: NUMBER OF SEQUENCES: 88
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
: STREET: 233 South Wacker Drive/6300 Sears Tower
: CITY: Chicago
: STATE: Illinois
: COUNTRY: United States of America
: ZIP: 60606-6402
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/474,379C
: FILING DATE: 07-JUN-1995
: CLASSIFICATION: 435
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/511,715
: FILING DATE: 20-APR-1990
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/206,188
: FILING DATE: 01-MAR-1994
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/688,352
: FILING DATE: 19-APR-1991
: ATTORNEY/AGENT INFORMATION:
: NAME: Clough, David W.
: REGISTRATION NUMBER: 36,107
: REFERENCE/DOCKET NUMBER: 27866/32771

```

```

: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (312) 474-6300
: TELEFAX: (312) 474-0448
: INFORMATION FOR SEQ ID NO: 13:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1721 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: CDNA
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 66..1274
: US-08-474-379C-13

Query Match 0.5%; Score 16; DB 2; Length 1721;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2731 cacagccgcgggccag 2746
DB 266 CACAGCCGCGGCCAG 281

RESULT 93
: US-09-146-249A-13
: Sequence 13, Application US/09146249A
: Patent No. 6069240
: GENERAL INFORMATION:
: APPLICANT: Wigler, Michael H.
: APPLICANT: Colicelli, John J.
: TITLE OF INVENTION: Cloning by Complementation and Related
: TITLE OF INVENTION: Processes
: NUMBER OF SEQUENCES: 85
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
: STREET: 6300 Sears Tower, 233 South Wacker Drive
: CITY: Chicago
: STATE: Illinois
: COUNTRY: United States of America
: ZIP: 60606-6402
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/146,249A
: FILING DATE:
: CLASSIFICATION:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/511,715
: FILING DATE: 20-APR-1990
: ATTORNEY/AGENT INFORMATION:
: NAME: Clough, David W.
: REGISTRATION NUMBER: 36,107
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 312/474-6300
: TELEFAX: 312-474-0448
: TELEX: 25-3856
: INFORMATION FOR SEQ ID NO: 13:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1721 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: CDNA
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 66..1274
: US-09-146-249A-13

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Query Match          0.5%; Score 16; DB 3; Length 1721;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

>Y 2731 cacagccgcgggccag 2746
>D 266 CACAGCCGCGGCCAG 281

RESULT 94
>S-08-206-188B-13
>Sequence 13, Application US/08206188B
>Patent No. 6100025
>GENERAL INFORMATION:
>APPLICANT: Wigler, Michael H.
>TITLE OF INVENTION: Cloning by Complementation and Related
>PROCESS
>NUMBER OF SEQUENCES: 84
>CORRESPONDENCE ADDRESS:
>ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
>STREET: 6300 Sears Tower, 233 South Wacker Drive
>CITY: Chicago
>STATE: Illinois
>COUNTRY: United States of America
>ZIP: 60606-6402
>COMPUTER READABLE FORM:
>MEDIUM TYPE: Floppy disk
>COMPUTER: IBM PC compatible
>OPERATING SYSTEM: PC-DOS/MS-DOS
>SOFTWARE: PatentIn Release #1.0, Version #1.25
>CURRENT APPLICATION DATA:
>APPLICATION NUMBER: US/08/206,188B
>FILING DATE: 01-MAR-1994
>CLASSIFICATION: 435
>PRIOR APPLICATION NUMBER: US 07/511,715
>FILING DATE: 20-APR-1990
>ATTORNEY/AGENT INFORMATION:
>NAME: Clough, David W.
>REGISTRATION NUMBER: 36107
>TELECOMMUNICATION INFORMATION:
>TELEPHONE: 312/474-6300
>TELEFAX: 312-474-0448
>TELEX: 25-3856
>INFORMATION FOR SEQ ID NO: 13:
>SEQUENCE CHARACTERISTICS:
>LENGTH: 1721 base pairs
>TYPE: nucleic acid
>STRANDEDNESS: single
>TOPOLOGY: linear
>MOLECULE TYPE: cDNA
>FEATURE:
>NAME/KEY: CDS
>LOCATION: 66..1274
>S-08-206-188B-13

Query Match          0.5%; Score 16; DB 3; Length 1721;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

>Y 2731 cacagccgcgggccag 2746
>D 266 CACAGCCGCGGCCAG 281

RESULT 95
>P-US91-02714-13
>Sequence 13, Application PC/TUS9102714
>GENERAL INFORMATION:
>APPLICANT: Wigler, Michael H.
```

```
>APPLICANT: Colicelli, John J.
>TITLE OF INVENTION: Cloning by Complementation and Related
>PROCESS
>NUMBER OF SEQUENCES: 55
>CORRESPONDENCE ADDRESS:
>ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
>Bicknell
>STREET: Two First National Plaza, 20 South Clark
>STREET: Street
>CITY: Chicago
>STATE: Illinois
>COUNTRY: USA
>ZIP: 60603
>COMPUTER READABLE FORM:
>MEDIUM TYPE: Floppy disk
>COMPUTER: IBM PC compatible
>OPERATING SYSTEM: PC-DOS/MS-DOS
>SOFTWARE: PatentIn Release #1.0, Version #1.25
>CURRENT APPLICATION DATA:
>APPLICATION NUMBER: PCT/US91/02714
>FILING DATE: 19910419
>CLASSIFICATION: 435
>PRIOR APPLICATION DATA:
>APPLICATION NUMBER: US 07/511,715
>FILING DATE: 20-APR-1990
>ATTORNEY/AGENT INFORMATION:
>NAME: Borun, Michael F.
>REGISTRATION NUMBER: 25447
>REFERENCE/DOCKET NUMBER: 27805/30197
>TELECOMMUNICATION INFORMATION:
>TELEPHONE: (312) 346-5750
>TELEFAX: (312) 984-9740
>TELEX: 25-3856
>INFORMATION FOR SEQ ID NO: 13:
>SEQUENCE CHARACTERISTICS:
>LENGTH: 1721 base pairs
>TYPE: NUCLEIC ACID
>STRANDEDNESS: single
>TOPOLOGY: linear
>MOLECULE TYPE: cDNA
>FEATURE:
>NAME/KEY: CDS
>LOCATION: 60..1274
>PCT-US91-02714-13

Query Match          0.5%; Score 16; DB 4; Length 1721;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2731 cacagccgcgggccag 2746
DB 266 CACAGCCGCGGCCAG 281

RESULT 96
US-08-481-814A-2
>Sequence 2, Application US/08481814A
>Patent No. 5869040
>GENERAL INFORMATION:
>APPLICANT: Hsu, Yen-Ming
>TITLE OF INVENTION: GENE THERAPY METHODS AND COMPOSITIONS
>NUMBER OF SEQUENCES: 12
>CORRESPONDENCE ADDRESS:
>ADDRESSEE: Biogen, Inc.
>STREET: 14 Cambridge Center
>CITY: Cambridge
>STATE: Massachusetts
>COUNTRY: United States of America
>ZIP: 02142
>COMPUTER READABLE FORM:
>MEDIUM TYPE: Floppy disk
>COMPUTER: IBM PC compatible
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OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/481.814A
FILING DATE:
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Kaplan, Warren A.
REFERENCE/DOCKET NUMBER: A001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-679-2000
TELEFAX: 617-679-2838
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1766 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cDNA to mRNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Homo sapiens
CELL LINE: HeLa
FEATURE:
NAME/KEY: mat_peptide
LOCATION: 429..1739
OTHER INFORMATION: /product= "E2F-2"
S-08-481-814A-2

Query Match 0.5%; Score 16; DB 2; Length 1766;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Y 2735 gccgcggccaggagg 2750
|||||

25 262 GCCGCGGCCAGGAGG 277

RESULT 97
US-08-993-228-1
Sequence 1, Application US/08993228
Patent No. 5976838
GENERAL INFORMATION:
APPLICANT: Jacobs, Kenneth
APPLICANT: McCoy, John M.
APPLICANT: Lavalie, Edward R.
APPLICANT: Racie, Lisa A.
APPLICANT: Merberg, David
APPLICANT: Treacy, Maurice
APPLICANT: Spaulding, Vikki
APPLICANT: Agostino, Michael J.
TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
TITLE OF INVENTION: ENCODING THEM
NUMBER OF SEQUENCES: 33
CORRESPONDENCE ADDRESS:
ADDRESSEE: Genetics Institute, Inc.
STREET: 87 Bridgepark Drive
CITY: Cambridge
STATE: MA
COUNTRY: U.S.A.
ZIP: 02140
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/993,228
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:

NAME: Sprunger, Suzanne A.
REGISTRATION NUMBER: 41,323
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 498-8284
TELEFAX: (617) 876-5851
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1790 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-993-228-1

Query Match 0.5%; Score 16; DB 2; Length 1790;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1581 tctctgtactggact 1596
|||||

Db 1235 TCTCTGCTACTGGACT 1250

RESULT 98
US-09-231-529-2
Sequence 2, Application US/09231529
Patent No. 6096308
GENERAL INFORMATION:
APPLICANT: Lal, Preeti
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
APPLICANT: Corley, Neil C.
APPLICANT: Shah, Purvi
TITLE OF INVENTION: HUMAN PROTEIN KINASE AND KINASE INHIBITORS
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Dr.
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/231,529
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/977,816
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0429 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-855-0555
TELEFAX: 650-845-4166
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1977 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: KIDNOT25
CLONE: 3453694
US-09-231-529-2

Query Match 0.5%; Score 16; DB 3; Length 1977;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1462 cagccccagcagagaa 1477

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419 CAGCCCCAGCAGAGAA 434

RESULT 99

US-09-255-911-1

Sequence 1, Application US/09255911

Patent No. 6013522

GENERAL INFORMATION:

APPLICANT: Brett P. Monia

APPLICANT: Lex M. Cowser

TITLE OF INVENTION: ANTISENSE MODULATION OF SMAD1 EXPRESSION

FILE REFERENCE: RTS-0040

CURRENT APPLICATION NUMBER: US/09/255,911

CURRENT FILING DATE: 1999-02-23

NUMBER OF SEQ ID NOS: 46

SEQ ID NO 1

LENGTH: 1990

TYPE: DNA

ORGANISM: Homo sapiens

FEATURE:

NAME/KEY: CDS

LOCATION: (433)..(1830)

US-09-255-911-1

Query Match

Best Local Similarity 0.5%; Score 16; DB 3; Length 1990;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1077 atggcccccagcatctg 1092

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1831 atggcccccagcatctg 1846

RESULT 100

US-08-619-554-7

Sequence 7, Application US/08619554

Patent No. 5821353

GENERAL INFORMATION:

APPLICANT: DOUGLAS, Cameron M.

APPLICANT: CHREBET, Gary L.

APPLICANT: CLEMAS, Joseph

APPLICANT: EL-SHERBEINI, Mohammed

APPLICANT: FOOR, Forrest

APPLICANT: KAHN, Jennifer

APPLICANT: KELLY, Rosemarie, - PARENT, S.A.

APPLICANT: MARRINAN, Jean, - RAMADAN, N.M.

APPLICANT: MORIN, Nancy, - REGISTER, E.A.

APPLICANT: ONISHI, Janet, - SHEI, Gan-Ju

TITLE OF INVENTION: DNA ENCODING 1,3 BETA-D GLUCAN

TITLE OF INVENTION: SYNTHASE SUBUNITS

NUMBER OF SEQUENCES: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: JOSEPH A. COPPOLA - MERCK & CO., INC.

STREET: 126 EAST LINCOLN AVENUE - P.O. BOX 2000

CITY: RAHWAY

STATE: NJ

COUNTRY: USA

ZIP: 07065

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/619,554

FILING DATE: 01-AUG-1996
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: COPPOLA, JOSEPH A
REGISTRATION NUMBER: 38,413
REFERENCE/DOCKET NUMBER: 19104PI
TELECOMMUNICATION INFORMATION:
TELEPHONE: 732-594-6734
TELEFAX: 732-594-4720
TELEX:

INFORMATION FOR SEQ ID NO: 7:

SEQUENCE CHARACTERISTICS:

LENGTH: 2069 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

US-08-619-554-7

Query Match

Best Local Similarity 0.5%; Score 16; DB 1; Length 2069;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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|||||

Db 1298 AGATTTTGGCTGAAGA 1313

Search completed: February 18, 2001, 14:23:44

Job time: 24889 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

.M nucleic - nucleic search, using sw model

Run on: February 18, 2001, 05:22:53 ; Search time 2148.29 Seconds
(without alignments)
9648.673 Million cell updates/sec

Title: US-09-434-382-3
Perfect score: 2958
Sequence: 1 ccggcgctaggtgaccggc.....aataaagattgagttgcaa 2958

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0
Searched: 7991742 seqs, 3503743858 residues

Word size : 0
Total number of hits satisfying chosen parameters: 15983484

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 100 summaries

Database :

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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	3	590	19.9	664	107	BE383336	BE383336 601298249
	4	590	19.9	692	107	BE382353	BE382353 601298656
	5	537	18.2	627	107	BE386924	BE386924 601274815
	6	528	17.8	761	135	BE795820	BE795820 601590856
	7	521	17.6	688	135	BE794311	BE794311 601591442
c	8	478	16.2	531	92	AW572950	AW572950 hf17h05.x
	9	465	15.7	823	110	BE619259	BE619259 601473130
	10	458	15.5	531	4	AA243700	AA243700 zr68g08.s
	11	451	15.2	451	17	AI200296	AI200296 qf86b12.x
	12	450	15.2	812	136	BE867512	BE867512 601443010
	13	448	15.1	499	91	AW510825	AW510825 hd40b11.x
	14	447	15.1	938	106	BE260626	BE260626 601146116
	15	442	14.9	493	92	AW575677	AW575677 UI-HF-BM0
	16	438	14.8	612	106	BE304720	BE304720 601106236
	17	436	14.7	493	7	AA455121	AA455121 zx78c04.s
c	18	425	14.4	478	92	AW592601	AW592601 hf45a09.x
	19	425	14.4	527	16	AI089646	AI089646 qb16g07.x
	20	421	14.2	421	20	AI468143	AI468143 tf92g05.x
	21	421	14.2	536	87	AW206103	AW206103 UI-H-B1-
	22	404	13.7	404	88	AW304130	AW304130 xs13e05.x
	23	402	13.6	474	4	AA291670	AA291670 zt37d04.s
	24	400	13.5	992	135	BE747163	BE747163 601577254
	25	384	13.0	574	105	BE250309	BE250309 600943455
	26	380	12.8	431	5	AA310236	AA310236 EST181085
	27	373	12.6	949	135	BE744197	BE744197 601577168
	28	370	12.5	370	137	BE883616	BE883616 601508091
	29	368	12.4	745	137	BE900936	BE900936 601674206
	30	352	11.9	518	144	R87541	R87541 ym89b04.r1
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	33	343	11.6	396	12	AA811170	AA811170 ob42c03.s
	34	342	11.6	446	10	AA634909	AA634909 ab27h02.r
	35	342	11.6	452	10	AA679618	AA679618 ag72c12.s
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	37	338	11.4	345	90	AW407520	AW407520 UI-HF-BM0
	38	334	11.3	677	135	BE742908	BE742908 601574609
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	42	323	10.9	397	10	AA632118	AA632118 np66h03.s
	43	323	10.9	482	144	R55841	R55841 yg89d01.r1
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	46	302	10.2	491	10	AA676661	AA676661 zj67h01.s
	47	301	10.2	480	144	R90875	R90875 ym10c02.r1
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60 276 9.3 461 10 AA635046 ab48b06.r
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62 273 9.2 457 14 AA994126 ou38b06.s
63 266 9.0 984 135 BE744876 601576324
64 255 8.6 410 15 AI033108 ow98908.s
65 255 8.6 461 92 AW592223 hf41a01.x
66 253 8.6 479 11 AA716607 s
67 250 8.5 501 5 AA311855 EST182568
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69 247 8.4 517 27 AI991599 ws18c04.x
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71 245 8.3 429 12 AA838624 oe91f04.s
72 243 8.2 346 96 AW889463 RC6-NT002
73 238 8.0 414 141 H22087
74 236 8.0 691 107 BE409312
75 235 7.9 439 17 AI201492 qs74b03.x
76 233 7.9 233 14 AF188525
77 233 7.9 394 141 H14462
78 232 7.8 865 8 AA522537 n138e08.s
79 228 7.7 282 13 AA928608 om75b03.s
80 228 7.7 282 145 T34024
81 227 7.7 433 19 AI357786
82 224 7.6 872 110 BE615669
83 221 7.5 291 4 AA235532
84 220 7.4 477 25 AI804749 tu42d02.x
85 218 7.4 228 88 AW296524
86 218 7.4 448 142 N36229
87 214 7.2 316 91 AW511765 xu76f03.x
88 211 7.1 249 88 AW247380 2820640.5
89 209 7.1 650 89 AW378247 RC1-HT021
90 207 7.0 219 8 AA504146 aa59e06.s
91 206 7.0 422 146 W37591
92 199 6.7 472 144 R51138
93 198 6.7 416 16 AI141263
94 193 6.5 394 6 AA346268
95 184 6.2 276 6 AA378232
96 182 6.2 290 147 Z44544
97 182 6.2 424 141 H03318
98 180 6.1 482 15 AI033342
99 179 6.1 577 110 BE619874
100 178 6.0 376 145 T72963 yc65b06.r1

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ALIGNMENTS

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RESULT 1
LOCUS BE260495 676 bp mRNA EST 13-JUL-2000
DEFINITION 601150702F1 NIH_MGC_19 Homo sapiens cDNA clone IMAGE:3503184 5',
mRNA sequence.
ACCESSION BE260495
VERSION BE260495.1 GI:9131807
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LLCM176 row: d column: 01
High quality sequence stop: 672.

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FEATURES             Location/Qualifiers
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                     /db_xref="taxon:9606"
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                     /clone_lib="NIH_MGC_19"
                     /tissue_type="neuroblastoma"
                     /lab_host="DH10B (phage-resistant)"
                     /note="Organ: brain; Vector: pOTB7; Site_1: XhoI; Site_2:
                     EcoRI; cDNA made by oligo-dr priming. Directionally
                     cloned into EcoRI/XhoI sites using the following 5'
                     adaptor: GGCAGCAG(G). Library constructed by Ling Hong
                     in the laboratory of Gerald M. Rubin (University of
                     California, Berkeley) using ZAP-cDNA synthesis kit
                     (Stratagene) and Superscript II RT (Life Technologies).
                     Note: this is a NIH_MGC Library."
BASE COORDINATES 154 a 207 c 176 g 139 t
ORIGIN
Query Match      22.9%; Score 676; DB 106; Length 676;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 676; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1143 acccagcattgtcctgaatgagaactgtgctcagttcacaaaccttcgcagcagccacaag 1202
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DB 1 ACCCAGCATTGTGCTCTGAATGAGAACTGTGCTCAGTTCACAACTTCGAGCCACACAG 60
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DB 61 ATTCAAAACCCAGCTCAACCTCATCCACCGGACATCTTCCCTCCTGCTCACCAGTTTCCGC 120
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QY 1323 taccagctcgtccccagagaggagtgagcagaggatgcccattattacttgcaatcctgag 1382
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DB 181 TACCAGCTCCGTCACAGAGGGAGTGGCAGAGGGATGCCATTATTACTTGCATCTGTAG 240
      |||||
QY 1383 gaattcatagttgaggcgtcagcttcccacttccacagcagagcgtgcagaggtacagag 1442
      |||||
DB 241 GAATTCATAGTTGAGGCGCTGCAGCTTCCCAACTTCCAGCAGAGCGTGCGAGGATACAG 300
      |||||
QY 1443 aggagtggcagagagggccccagccccccagcagagagagaaagagtcagtcaccagaatcac 1502
      |||||
DB 301 AGGAGTGGCGCAGGAGCGGCCCGCCAGCAGAGAGAGAAAGTCAAGTACCAAGAAATCATC 360
      |||||
QY 1503 ttccctggaacagggtctgctccatcccgatgaagattcgaatgtcagtcagtcaccactgtc 1562
      |||||
DB 361 TTCTTTGGNACAGGGTCTGCCATCCCGATGAGATTCGAAATGTCAGTGCACACACTTGTG 420
      |||||
QY 1563 aacataagccccgcagacgtctctgtactgtgactgtggtgagggcacattttggggcagctg 1622
      |||||
DB 421 AACATAAGCCCGCACACAGCTCTCTGCTACTGTGACTGTGTGTGAGGGACATTTGGGACGCTG 480
      |||||
QY 1623 tccgtcattacggagagaccaggtgacagaggttcctctgggacacctggctgtgtttgtg 1682
      |||||
DB 481 TCCCGTCAATTACGAGAGACCAGGTGACAGGGTCTCTGGGACCCCTGCTGTGTTTGTG 540
      |||||
QY 1683 tccccacctgcagcagatcacacacagggcttgcgaagtattcttgcgcagagagaacgc 1742
      |||||
DB 541 TCCCACCTGCACGCAGATCACACACAGGGCTTGGCCAAAGTATCTTGTGTCAGAGAGAACGC 600
      |||||
QY 1743 gctttggcatctttgggaaagccgttccaccttctgtgtgtgtgtgtgtgtgtgtgtgtgt 1802
      |||||
DB 601 GCCTTGGCATCTTTGGGAAAGCCGCTTACCCTTTGCTGGTGTGCTGCCCAACCAAGCTC 660
      |||||
QY 1803 aaagcctggctccagc 1818
      |||||
DB 661 AAAGCCTGGCTCCAGC 676

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RESULT 2
LOCUS BE250412/c 975 bp mRNA EST 13-JUL-2000
DEFINITION 600943455T1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:2960077 3',
mRNA sequence.
ACCESSION BE250412
VERSION BE250412.1 GI:9120523
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 975)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LLCM52 row: n column: 14
High quality sequence start: 22
High quality sequence stop: 756.
FEATURES
Location/Qualifiers
1..975
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2960077"
/tissue_type="rhabdomyosarcoma"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: muscle; Vector: pOTB7; Site:1: EcoRI;
Site:2: XhoI; CDNA made by oligo-dT priming.
Directionally cloned into EcoRI/XhoI sites using the
following 5' adaptor: GGCACGAG(G). Size-selected >500bp
for average insert size 1.8kb. Library constructed by
Ling Hong in the laboratory of Gerald M. Rubin (University
of California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT 185 a 296 c 258 g 236 t
ORIGIN
Query Match 20.8%; Score 614; DB 105; Length 975;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 614; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
2197 acgaggagttcattatgtgaacacattcagcagcgctatgccaaggtccccctcttca 2256
711 ACGGGAGTTCAATTATGCTCAACCACTTACGACGCGCTATGCAAGGTCCCTCTTCA 652
2257 gccccaacttcagcagaaagtgggaagttgctctttgacacatgaaggtctgttggag 2316
651 GCCCAACTTTCAGCAGAAAGTGGAGTTCCTTTTGACCATGAAAGTCTGCTTTGGAG 592
2317 actttcaaatgccaaactatccccactgaagccctgtttgtgggacatcg 2376
591 ACTTTCACAAATGCCCAAGCTATTCCCTCCACTGAAAGCCCTGTTTGCCTGGGACATCG 532
2377 agagatggaggagcgcagggagaagcggagctcgccgaggtgcggcgccctctctgt 2436
531 AGGAGATGGAGGAGCGCAGGAGAGCGGAGTGGCGAGGTGCGGCGGCCCTCTCTGT 472
2437 ccagggagctggcagcgcgctggagatgggagcctcagcagaagcgggcccacacag 2496
471 CCAGGGAGCTGGCAGGGCGGCTGGAGATGGGAGCTCAGCAGAAAGCGGGCCACACAG 412
2497 aggagccacagggccaagaaggtcagagcccgagtgaaagtctgggagaccctgaactcga 2556

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Db 411 AGGAGCCACAGGCCAAGAGGTGAGAGCCAGTGGAGACCCCTGAACTCAGA 352
QY 2557 agcgtgtgttcttctgtcccccacgcacgcacgcgtatctgcctctgtgtgtagaagc 2616
Db 351 AGGCTGTGTCTCTCTGCCCCACGACGACCGATCTGCCCTCCCTTGTGTGTAAGC 292
QY 2617 tgaagagacggtccccccagggagcagctcaggatagggtggtatggagctgtccaggc 2676
Db 291 TGAAGACACGGTCCCGCCAGGAGGCGCTAGGATAGGTGATGGAGCTGCCGAGGC 232
QY 2677 ttgggtccacataagcactagctctatagatgcctcttaggactgtgctgtgcgcacgc 2736
Db 231 TTGGGCTCCCATAGACACTAGTCTATAGATGCCTCTTAGGACTGGTGGCTGGCACAGC 172
QY 2737 cgcggccagggaggtgcccacgcagcagcagcagcagcagcagcagcagcagcagcagcag 2796
Db 171 CGCGGCGCAGGAGGCTGCCACACGGAAGCAGATGAACCTAATTTCAATTTCAAGGCAG 112
QY 2797 tttttaagaagtc 2810
Db 111 TTTTAAAGAGTC 98
RESULT 3
LOCUS BE383336 664 bp mRNA EST 21-JUL-2000
DEFINITION 601298249F1 NIH_MGC_19 Homo sapiens cDNA clone IMAGE:3628308 5',
mRNA sequence.
ACCESSION BE383336
VERSION BE383336.1 GI:9328701
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 664)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LLCM313 row: a column: 13
High quality sequence stop: 662.
FEATURES
Location/Qualifiers
1..664
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3628308"
/tissue_type="neuroblastoma"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: brain; Vector: pOTB7; Site:1: XhoI; Site:2:
EcoRI; CDNA made by oligo-dT priming.
Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Library constructed by Ling Hong
in the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
Note: this is a NIH_MGC library."
BASE COUNT 154 a 199 c 174 g 137 t
ORIGIN
Query Match 19.9%; Score 590; DB 107; Length 664;
Best Local Similarity 100.0%; Pred. No. 0;

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JOURNAL COMMENT
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 Tel: (301) 496-1550
 Email: Robert_Strausberg@nih.gov
 This clone is available royalty-free through LLNL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Seq primer: -40UP from Gibco
 High quality sequence stop: 459.
 Location/Qualifiers
 1..531
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:2932185"
 /clone_lib="Soares_NFL_T_GBC_S1"
 /lab_host="DH10B"
 /note="Organ: pooled; Vector: pT73D-Pac (Pharmacia) with
 a modified polylinker; Site_1: Not I; Site_2: Eco RI;
 Equal amounts of plasmid DNA from three normalized
 libraries (fetal lung NCHL19W, testis NHT, and B-cell
 NCI-CGAP-GCB1) were mixed, and ss circles were made in
 vitro. Following HAP purification, this DNA was used as
 tracer in a subtractive hybridization reaction. The driver
 was PCR-amplified cDNAs from pools of 5,000 clones made
 from the same 3 libraries. The pools consisted of
 I.M.A.G.E. clones 297480-302087, 682632-687239,
 726408-728711, and 729096-731399. Subtraction by Bento
 Soares and M. Fatima Bonaldo."
 99 a 154 c 136 g 142 t

BASE COUNT
 ORIGIN
 Query Match 16.2%; Score 478; DB 92; Length 531;
 Best Local Similarity 99.8%; Pred. No. 1.3e-241;
 Matches 528; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 2430 ccctgtccaggagctggtgagcggcctgagagtgaggagccctcagcagagcgggcc 2489
 531 CTCCTGTCCAGGAGCTGCAGCGCGCCTGGAGATGGGAGCCCTCAGCAGAGCGGGCC 472
 2490 cacacagaggagccagcagcgaaggtcagagccagtcagagtcctgggagacccctga 2549
 471 CACACAGAGGAGCCACAGGCCAAGAAGGTGAGAGCCAGTGAAGATCTGGAGACCTGA 412
 2550 actcagaggtgtgtgtcttctgcccacgacgacccgctatctgcccctctgctgg 2609
 411 ACTCAGAGGCTGTGTGTCTTCTGCCCCACGACGACCCGATCTGCCCTCTTCTGCTGG 352
 2610 tagaagctgaagagcacggtcccccagagggcagctcagagtaggtgtgtagagctgtg 2669
 351 TAGAAGCTGAAGAGCACGCTCCCCAGGAGGAGCTCAGAGTAGGTGTATGGAGCTGTG 292
 2670 ccgaggtgtgggtccacataagcactagtctatagatgcctcttaggactggtgctgt 2729
 291 CCGAGGCTTGGGCTCCACATAGCACTAGTCTATAGATGCCCTCTTAGACTGGTGCCTG 232
 2730 gcacagcgcggggcaggaggtgcccacaggaagcagacagataaactaattcatttc 2789
 231 GCACAGCCCGGGCCAGGAGGTGCCACAGGAGCAAGCATGAATTAATTTCAATTC 172
 2790 aaggcagtttttaagaagctcttggaacagagcgggcaccttctccttaataccagcaa 2849
 171 AAGGAGTTTTTAAGAAGTCTATGGAACAGAGCGGGCACCCTCTTCTATATCCAGCAA 112
 2850 agtgattccctgcacaccagagacagcagagtaaacagatcagtggtgtcctaagtctcg 2909
 111 AGTGATTCCTGCACACCAGAGCAAGCAGAGTAAACAGATCAGTGGGTCTAAGTGTCCG 52
 2910 acacttaacgaaatagatttcacgtgcaataaagattgattgcaa 2958
 51 AGACTTAACGAAATAGTATTTCAGCTGCAATAAAGATTGATTGTGCAA 3
 RESULT 9

BE619259 823 bp mRNA EST 24-AUG-2000
 LOCUS 601473130F1 NIH_MGC_68 Homo sapiens cDNA clone IMAGE:3876223 5',
 DEFINITION mRNA sequence.
 ACCESSION BE619259
 VERSION BE619259.1 GI:9890197
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 823)
 NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/
 National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished (1999)
 JOURNAL Contact: Robert Strausberg, Ph.D.
 COMMENT Tel: (301) 496-1550
 Email: Robert_Strausberg@nih.gov
 Tissue Procurement: DCTD/DTP/Gazdar
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)
 Clone Distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LICM623 row: k column: 08
 High quality sequence stop: 695.
 Location/Qualifiers
 1..823
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:3876223"
 /clone_lib="NIH_MGC_68"
 /tissue_type="large cell carcinoma"
 /lab_host="DH10B (phage-resistant)"
 /note="Organ: lung; Vector: pCMV-SPORT6; Site_1: NotI;
 Site_2: SalI; Cloned unidirectionally. Primer: Oligo dI.
 Average insert size 1.8 kb. Library constructed by Life
 Technologies."
 185 a 219 c 259 g 160 t

BASE COUNT
 ORIGIN
 Query Match 15.7%; Score 465; DB 110; Length 823;
 Best Local Similarity 100.0%; Pred. No. 1e-234;
 Matches 465; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 1678 ttgttccacacctgcacgcagatcacacacgggcttcccaagtatcttgcgcagag 1737
 75 TTGTGTCACCTGCACGCAGATCACACACGGGCTTCCCAAGTATCTGCTGCAGAGAG 134
 1738 aacgcgcttggcattcttgggaaagccgctcacccttgcctggtggtgccccacc 1797
 135 AACGCGCTTGGCATCTTTGGGAAAGCGCTTACCCCTTGTGTTGTTGCCCAACC 194
 1798 agctcaagcctggtccagcagcagcagcagcagcagcagcagcagcagcagcagc 1857
 195 AGCTCAAGCCTGGCTCCAGCAGTACCAACACAGTCCAGAGGTCTCTGCACCATCA 254
 1858 gtatgattccctggcaaatgccttcaggaggggctgagatctccagtcctcagtgga 1917
 255 GTATGATTCTCCCAATGCCTTCCAGGAAGGGCTGAGATCTCCAGTCTCCAGTGGAAA 314
 1918 gattgatcagtcgctgttgcgaacatgtattgggaagatttcagaccctgtctggtgc 1977
 315 GATTGATCAGTTCGCTGTGGCAACATGTATTTGGAAGAGTTCAGACCTGTCTGTGTC 374
 1978 ggcactgcaacatgcgttgcgtgcgtgcgtgcgtgcgtgcgtgcgtgcgtgcgtgc 2037
 375 GCACCTGCAAGCATCGCTTGGCTGCGCTGGTGCACACCTCTGCTGGAAGTGGTCT 434
 2038 attccgggggacacatgcccctgcaggctcctggtcggatgggaaagatgccaccctcc 2097

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b 435 ATTCCGGGACACCATGCCCTCGAGGCTCTGGTCGGATGGGAAGATGCCACCTCC 494
y 2098 tgatacatgaagccaccctcggaagatggttggaagagaagcag 2142
b 495 TGATCATGAAGCCACCCCTGGAAGATGTTTGAAGAGAGCAG 539

RESULT 10
AA243700 531 bp mRNA EST 07-MAR-1997
DEFINITION zr68g08.s1 Soares_NHMPu_S1 Homo sapiens cDNA clone IMAGE:668606 3'
similar to SW:YK59_YEAST P36159 HYPOTHETICAL 96.8 KD PROTEIN IN
SI52-MTD1 INTERGENIC REGION. ; mRNA sequence.
ACCESSION AA243700
VERSION AA243700.1 GI:1874492
SYNOPSIS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 531)
AUTHORS Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
,R., Williamson,A., Wohldmann,P. and Wilson,R.
TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.wustl.edu
This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Possible reversed clone: similarity on wrong strand
Seq primer: -41m13 fwd. Et from Amersham
High quality sequence stop: 466.
Location/Qualifiers
1. 531
/organism="Homo sapiens"
/db_xref="GDB:5562573"
/db_xref="taxon:9606"
/clone="IMAGE:668606"
/clone_lib="Soares_NHMPu_S1"
/tissue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/lab_host="DH10B"
/note="Organ: mixed (see below); Vector: pT73D-Pac
(Pharmacia) with a modified polylinker; Site_1: Not I;
Site_2: Eco RI; Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NbHM, pregnant uterus
NbHPU, and fetal heart NBH19W) were mixed, and ss circles
were made in vitro. Following HAP purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of I.M.A.G.E. clones 260232-265223,
340488-345479, and 484488-489479."
BASE COUNT 144 a 137 c 143 g 107 t
ORIGIN
Query Match 15.5%; Score 458; DB 4; Length 531;
Best Local Similarity 100.0%; Pred. No. 5e-231;
Matches 458; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
y 251 ccggagctcgggcgccgctctacgtctctccgagttcaaccggtatctcttcaactg 310
b 74 CCGGACTCGGGCGCGCTACGTCCTCCGAGTTCACCGGTACTCTCTCAACG 133
y 311 tggagaagcggttcagagactatcaggagcacaagttaaagggttcgctcgccggacaa 370

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Db 134 TGGAGAAGCGGTTGAGACACTCATCGAGGACACAAAGTTAAAGGTTGCTCGCCTGGACAA 193
Qy 371 catattctgcacacgaatgcactgctcttaagtgtggggcttaagtgaatgattctttac 430
Db 194 CATATTCTGCACACGAATGCACCTGGTCTAAATGTTGGGGCTTAAAGTGAATGATTCTTAC 253
Qy 431 tttaaaggaaccggggttcctcaaatgtgtacttcttgagacctccacaactgagaaata 490
Db 254 TTAAAGGAACCCGGCTTCCAAAGTGTACTTCTTGACCTCCACAACTCGAAAAATA 313
Qy 491 cctcgaagcaatcaaaaattttcttggtccattgaaaggaatagaaactggtgtgcggtc 550
Db 314 COTCGAAGCAATCAAAATATTTCTGGTCCATTGAAGGAATAGAATGGCTGTGGGGCC 373
Qy 551 ccactctgcccagaatcacgaggtgaaccatgacagtttaccagatccccatacacag 610
Db 374 CCACCTCTGCCCCAGAAATACGAGGATGAACACCATGACAGTTTACCAGATCCCATACACAG 433
Qy 611 tgaacagagggagggaagcaccacacatgagcagatccagagtcagaaaggcctctcagcggct 670
Db 434 TGAACAGAGGAGGGGAAAGACCAACCATGGCAGAGTCCAGAAAGGCCCTCTCAGCAGGCT 493
Qy 671 cagtcagagcgatcttcagactccgagtcgagtgcaatgaaa 708
Db 494 CAGTCCAGAGCGATCTTCAGACTCCGAGTCGGAATGAAA 531

RESULT 11
AI200296 451 bp mRNA EST 14-OCT-1998
LOCUS qf86b12.x1 Soares_fetal_lung_NbH19W Homo sapiens cDNA clone
DEFINITION IMAGE:1756895 3' similar to SW:YK59_YEAST P36159 HYPOTHETICAL 96.8
KD PROTEIN IN SI52-MTD1 INTERGENIC REGION. ; mRNA sequence.
ACCESSION AI200296
VERSION AI200296.1 GI:3752902
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 451)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -40UP from Gibco
High quality sequence stop: 442.
Location/Qualifiers
1. 451
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1756895"
/clone_lib="Soares_fetal_lung_NbH19W"
/dev_stage="19 weeks"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: lung; Vector: pT73D (Pharmacia) with a
modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer
[5'-TGTTACCAATCTGAAGTGGAGCGCGCAATTTTTTTTTTTT-3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pT73 vector
(Pharmacia). Library went through one round of
normalization to a Cot = 5. Library constructed by Bento
Soares and M.Fatima Bonaldo. This library was constructed
from the same fetus as the fetal heart library, Soares
fetal heart NBH19W."
FEATURES
Source

```


JOURNAL COMMENT
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -40UP from Gibco
High quality sequence stop: 470.

FEATURES
Location/Qualifiers
1..499
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2911965"
/clone_lib="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; Equal amounts of plasmid DNA from three normalized libraries (fetal lung NBHL19W, testis NHT, and B-cell NCI-CGAP-GCB1) were mixed, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 297480-302087, 682632-687239, 726408-728711, and 729096-731399. Subtraction by Bento Soares and M. Fatima Bonaldo."
94 a 142 c 126 g 137 t

BASE COUNT
ORIGIN
Query Match 15.1%; Score 448; DB 91; Length 499;
Best Local Similarity 99.8%; Pred. No. 1e-225;
Matches 498; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

2460 gaggatggggagcctcagcagagcggggccacacagagagccagccagcgaaggtc 2519
|||||
2499 GAGGATGGGGAGCCTCAGCAGAAGCGGGCCACACAGAGGAGCCACAGCCAAAGGTC 440
|||||
2520 agagccagtggaagatctggggagccctgaactcgaagcgtgtgtttctgcccac 2579
|||||
2439 AGAGCCAGTCAGAGATCTGGGAGACCTGAACCTCAGAGGCTGTGTCTTCTGCCCCAC 380
|||||
2580 gcacacacccgtatctccctcttctgtgtgtagaagctgaagcagcagctccccaggag 2639
|||||
2379 GCACGACCCGTCATCTGCCCTCTTCTGCTGTAAGCTGAAGAGCAGCGTCCCCAGGAG 320
|||||
2640 gcagctcagataggtggtatgagctgtgcgagcgttgggtctccacataagcactag 2699
|||||
2319 GCAGCTCAGATAGGTGGTATGAGCTGTGCGAGGCTTGGCTCCCATTAAGCACATAG 260
|||||
2700 tctatagatgcctcttagactgtgtcctggtgacagcgcgggaggggtgccaac 2759
|||||
2259 TCTATAGATGCTCTTAGACTGTGTGCTGTCACAGCCGCGGAGGAGGTGCCACAC 200
|||||
2760 ggaagcagcagatgaactaatctatttcaaggcagtttttaagaagatcttggaaaca 2819
|||||
2199 GGAAGCAAGCAGATGAACCTAATTTTCATTAAGGCGAGTTTAAAGAGTCATGGAACA 140
|||||
2820 gacgcgccaccttctccttaactcagcaaaagtattcctcctcacacagacagcag 2879
|||||
2139 GACGGGGGACCTTTCCTCTTANTCCAGCAAAAGTATTCCCTGCACACCCAGACAGCAG 80
|||||
2880 agtaacagatcagtggtgttaagtgtccgagacttaacgaaatagtatttcagtcca 2939
|||||
279 AGTAACAGATCAGTGGGTCCTAAGTGTCCGAGACTTAACGAAATAGTATTTCAGCTGA 20
|||||
2940 ataaagattgagttgcaa 2958
|||||
219 ATAAAGATTGAGTTTGCAA 1

RESULT 14

BE260626 938 bp mRNA EST 13-JUL-2000
LOCUS 601146116F1 NIH_MGC_19 Homo sapiens cDNA clone IMAGE:3161691 5',
DEFINITION mRNA sequence.
ACCESSION BE260626
VERSION BE260626.1 GI:9132065
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 938)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LLCml19 row: O column: 04
High quality sequence stop: 621.

FEATURES
Location/Qualifiers
1..938
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3161691"
/clone_lib="NIH_MGC_19"
/tissue_type="neuroblastoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: brain; Vector: pORF7; Site_1: XhoI; Site_2: EcoRI; CDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCAGGAG(G). Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC Library."
190 a 263 c 283 g 202 t

BASE COUNT
ORIGIN
Query Match 15.1%; Score 447; DB 106; Length 938;
Best Local Similarity 100.0%; Pred. No. 3.5e-225;
Matches 447; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 966 ttgtgtgtgtagaatgtccagatgaaagcttcattcaaccatctgtgagaatgccacc 1025
|||||
DB 135 TTTGTGGTGGTAGAATGTCAGATGAAAGCTTCATTCAACCCATCTGTGAGAAATGCCACC 194
|||||
QY 1026 tticagaggtaccagaaagcagatgccccgtggtgtgtgttcacatggcccca 1085
|||||
DB 195 TTTTCAGAGTACCAAGAAAGCAGATGCCCGCTGGCCCTTGGTTCACATGSCCCCA 254
|||||
QY 1086 gcatctgtctgtgtgagcagcaggtaccagcagtgatggagaggtttgggctgcacacc 1145
|||||
DB 255 GCATCTGTGCTTGTGGACAGCAGGTACCAGCAGTGGATGGAGAGGTTTGGGCTGCACACC 314
|||||
QY 1146 cagcacttggtcctgaatgagaactgtgcctcagttcacaaaccttcgagccacaagatt 1205
|||||
DB 315 CAGCACTTGGTCTCTGAATGAGAACTGTGCTCAGTTCACAACTTCGCGAGCCACAAGATT 374
|||||
QY 1206 caaacccagctcaacctcatccaccgcagacatcttccccctgctcaccagtttcgctgt 1265
|||||
DB 375 CAACCCAGCTCAACCTCATCCACCCGGACATCTTCCCCCTCTCACCAGTTTCGCTGT 434
|||||
QY 1266 aagaaggaggggccccaccctcagtggtgccatggttcaggggtgaatgcctcctcaagtac 1325
|||||
DB 435 AAGAGGAGGGGGCCCCACCCCTCAGTGTGCCCATGTTTCAGGGTGAATGCTCCTCCTCAAGTAC 494
|||||

```
1326 cagctcgtcccaaggaggagtgagcagaggtgagcattattacttgcaactcctgaggaa 1385
|||||
495 CAGCTCGTCCCGAGGAGGAGTGGCAGAGGATGCCATTATTCTTGAATCCTGAGGAA 554
|||||
1386 ttcatagtgagcgctcagcttccc 1412
|||||
555 TTCATAGTGGAGCGGTGCAGCTCCC 581
|||||

RESULT 15
LOCUS AW575677 493 bp mRNA EST. 15-MAR-2000
DEFINITION UI-HE-BM0-adl-b-07-0-UI.s1 NIH_MGC_38 Homo sapiens CDNA clone
IMAGE:3061957 3', mRNA sequence.
ACCESSION AW575677
VERSION AW575677
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 493)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Oligo-dr track not found. Not 1 site shown in beginning of sequence
is likely internal to the message. Tissue Procurement: Louis M.
Staudt, M.D., Ph.D.
CDNA Library Preparation: M.B. Soares Lab
CDNA Library Arrayed by: M.B. Soares Lab
DNA Sequencing by: M.B. Soares Lab
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bbrp/image/image.html
Seq primer: M13 Forward
POLYA=NO.

FEATURES
Location/Qualifiers
1..493
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3061957"
/clone_lib="NIH_MGC_38"
/tissue_type="lymph"
/cell_type="germinal center B cells"
/cell_line="MGC85"
/lab_host="DH10B (LT1)"
/notes="Vector: pT73-Pac; Site_1: NotI; Site_2: Eco RI;
Constructed from size fractionated cytoplasmic mRNA
(2.5-3.5kb). Directionally cloned. Cells provided by Louis
M. Staudt, Ph.D. Library preparation by Maria de Fatima
Bonaldo, Ph.D. and M. Bento Soares, Ph.D."
BASE COUNT 126 a 131 c 138 g 97 t
ORIGIN
|||||

Query Match 14.9%; Score 442; DB 92; Length 493;
Best Local Similarity 99.8%; Pred. No. 1.5e-222;
Matches 492; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

141 cggccgcgaagaccctgtcgccacctcgccacgcagagagcgcgacctcgggg 200
|||||
1 CGCCCGCGCAAGACCCGCTGCGCGACCTCGCCACGCGAGAGGCGCGACGCTCGGG 60
|||||
201 tgcctcgccggcccaaacacgcgtgtacctgaggtggtgagcggtgagcgactcg 260
|||||
61 TGTCTCGGGCGGCCCAACACCGTGTACCTGACAGTGTGTCACGCGGTGACCGGGACTCG 120
|||||
261 ggcgcgcgcgtctacgtcttctccaggttcaaccggtatctcttcaactgtggaagcc 320
|||||
```

```
Db 121 GGCGCGCGCTCTACGTCTTCTCCGAGTTCAACCGGTATCTCTTCAACTGTGGAGAAGGC 180
Qy 321 gttcagagactcatgagagagcacaaagtttaaggttgcctgcctgggacacacatattcctg 380
|||||
Db 181 GTTCAGAGACTCATGTCAGGAGCACAAGTTAAAGGTTGCTCGCTGGACACATATTCTCTG 240
Qy 381 acacgaatcacctggtcctaatgtttggggccttaagtggaatgattcttctttaaaggaa 440
|||||
Db 241 ACAGTAATGCACTGGTCTAATGTGGGGCTTAAGTGGGAATGATTCTTCTTTAAAGGAA 300
Qy 441 accgggtctccaaagtgtgtactttctgacctcccaacacacacacacacacacacacacac 500
|||||
Db 301 ACCGGGCTTCCAAAGGTGTACTTTCTGCACCTCCACAACTGCAAAATACCTCGAAGCA 360
Qy 501 atcaaaatattttctggtccattgaaagaatagaactggtgtgctggcggcccaactctgcc 560
|||||
Db 361 ATCAAAATATTTTCTGCTCCATTGAAAGGAATAGAACTGGCTGTGCGGNCCTACTCTGCC 420
Qy 561 ccagaatcacgagatgaaccatgacagttaccagatcccatcacacacacacacacacacag 620
|||||
Db 421 CCAGTAATAGAGATGAACCAATGACAGTTACAGATCCCATACACAGTGAACAGAGG 480
Qy 621 aggggaaagcacc 633
|||||
Db 481 AGGGGAAGCACC 493
|||||

RESULT 16
LOCUS BE304720 612 bp mRNA EST 13-JUL-2000
DEFINITION 601106236F1 NTH_MGC_15 Homo sapiens CDNA clone IMAGE:3349304 5',
mRNA sequence.
ACCESSION BE304720
VERSION BE304720.1 GI:9176150
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 612)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: NIH Intramural Sequencing Center
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: Image.llnl.gov
Plate: LILCM143 row: h column: 09
High quality sequence start: 21
High quality sequence stop: 609.
Location/Qualifiers
1..612
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3349304"
/clone_lib="NIH_MGC_15"
/tissue_type="adenocarcinoma cell line"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: colon; Vector: pOTB7; Site_1: XhoI; Site_2:
EcoRI; CDNA made by oligo-dr priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(S). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)"
BASE COUNT 143 a 178 c 161 g 130 t
ORIGIN
```

```
Query Match      14.8%; Score 438; DB 106; Length 612;
Best Local Similarity 100.0%; Pred. No. 2e-220;
Matches 438; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

>Y 1041 ggaagcagatgccccgctggtgtgttccatgagcccgagcattctgtctgtg 1100
>D 108 GGAAGGAGAGATGCCCCGCTGTGTGTGTTCATATGGCCGAGCATCTGTCTTGTG 167
>Y 1101 gacagcaggtaccagcagtgatgagaggtttggcctgacaccagcacttggtcctg 1160
>D 168 GACAGCAGGTACCAGCAGTGTGATGAGAGGTTTGGGCTGACACCCAGCAGCTTGTCCTG 227
>Y 1161 aatgagaactgtgctcagttcacaaccttgcagcccaagattcaaccagctcaac 1220
>D 228 AATGAGAACTGTGCTCAGTTTCAACACCTTCGACGCCCAAGATTCAACCCAGCTCAAC 287
>Y 1221 ctcatccaccggacatcttccccctgtccaccagtttcccgctgtaagaaggagggcccc 1280
>D 288 CTCATCCACCCGAGACATCTTCCCCCTGTCTACACGTTTCCGCTGTAAAGAGGGGCCCC 347
>Y 1281 accctcagtgcccatggttcaggtgtaagtgcctccctcaagttaccagctccgtcccccag 1340
>D 348 ACCCTCAGTGTGCCATGTTTCAAGGTTGAATGCCCTCTCAAGTACCAGCTCCGTCCTCCAGG 407
>Y 1341 agggagtgccagaggtgacattattcttgatcctcaggtgaggaattcagttgagcgc 1400
>D 408 AGGGAGTGGCAGAGGAGTGCCTATTATTCTTGAATCTCGAGGAATTCATAGTTGAGGCG 467
>Y 1401 ctgcagcttcccaactccacagagcgtgcagaggtacagaggtgcagagagcgcgcgc 1460
>D 468 CTGCAGCTTCCCACTCCAGCAGAGCGTGCAGAGGTACAGAGGAGTGCAGGACGCGC 527
>Y 1461 ccagcccccagcagagaaa 1478
>D 528 CCAGCCCCAGCAGAGAAA 545

RESULT 17
LOCUS      AA455121      493 bp      mRNA      EST      06-JUN-1997
DEFINITION zx78c04.sl Soares ovary tumor N6HOT Homo sapiens cDNA clone
            IMAGE:809862 3', similar to SW_YK59_YEAST P36159 HYPOTHETICAL 96.8
            KD PROTEIN IN SIS2-WD1 INTERGENIC REGION. ;, mRNA sequence.
ACCESSION  AA455121
VERSION     AA455121.1  GI:2177897
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
            Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
            Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B.,
            Schellenberg, K., Steptoe, M., Tan, F., Theising, B., White, Y., Wyllie
            , T., Waterston, R., and Wilson, R.
            WashU-Merck EST Project 1997
            Contact: Wilton RK
            Washington University School of Medicine
            444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: est@watson.wustl.edu
            This clone is available royalty-free through LNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            Possible reversed clone: similarity on wrong strand.
            Seq primer: -41m13 fwd. ET from Amersham
            High quality sequence stop: 447.
            Location/Qualifiers
            source          1. .493
```

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/organism="Homo sapiens"
/db_xref="CDB:6039680"
/db_xref="taxon:9606"
/clone="IMAGE:809862"
/clone_lib="Soares ovary tumor N6HOT"
/sex="Female"
/tissue_type="ovarian tumor"
/lab_host="DH10B (ampicillin resistant)"
/notes="Organ: ovary; Vector: p773D (Pharmacia) with a
modified polylinker; Site: 1: Not 1; Site 2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTTACCAATCTGAAGTGGAGGCGCGCGGTTTTTTTTTTTTTTT 3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified p773 vector
(Pharmacia). Library constructed by Bento Soares and
M.Fatima Bonaldo."
BASE COUNT      131 a      128 c      130 g      104 t
ORIGIN
Query Match      14.7%; Score 436; DB 7; Length 493;
Best Local Similarity 100.0%; Pred. No. 2.3e-219;
Matches 436; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

>Y 252 cgggactcggcgccgctcagctcttccaggttcaaccggtatctctcaactgt 311
>D 58 CGGGACTCGGCGCGCGCTCTACGTCTTCCGAGTTCAACCGGTATCTCTTCAACTGT 117
>Y 312 ggagaagcgttcagagactcatgcaggagcacaagttaaaggttgctgcctggacaac 371
>D 118 GGAGAGCGGTTTCAGAGACTCATGCAGGAGCACAAGTTAAAGGTGCTCGCTGGACAC 177
>Y 372 atattcctgacacgaatgcactggtctaatgttgggggttaagtggagtgattttact 431
>D 178 ATATTCTTGACACGAATGCAGTGGTCTAATGTGGGGGCTTAAGTGAATGATCTTACT 237
>Y 432 ttaaggaacacgggcttccaaagtgtactttctgacacctccacaactgaaaaaac 491
>D 238 TTAAGGAACACGGGCTTCCAAAGTGTGTACTTTCTGGACCTCCACAACTGGAAAAATAC 297
>Y 492 ctgaagcaatcaaaatatttctggttcattgaaaggaatagaactggtgtcgggccc 551
>D 298 CTCGAAGCAATCAAAATATTTCGTCTCATTTGAAGGAATAGAACTGGCTGTGGGCC 357
>Y 552 cactgtccccagaatacagagatgaaccatgcagctttaccagatccccatcacagt 611
>D 358 CACTCTGCCCCAGAAATACGAGATGAAACCATGACAGTTTACAGATCCCCATACACAGT 417
>Y 612 gaacagagaggggaaagcaccacacccatggcagagtcagaaaggccctcagaggctc 671
>D 418 GAACAGAGAGGGGAAAGCACCACCAACCATGGCAGAGTCCAGAAAGGCCCTCTCAGCAGGCTC 477
>Y 672 agtcacagagcgtctt 687
>D 478 AGTCGAGAGCGGATCTT 493

RESULT 18
LOCUS      AW592601/c      478 bp      mRNA      EST      22-MAR-2000
DEFINITION hf45a09.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
            IMAGE:2934808 3', mRNA sequence.
ACCESSION  AW592601
VERSION     AW592601.1  GI:7279786
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
            NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
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```
299  CACCTTTCAGAGTACACAGGAAGGACAGATGCCCGCTGGCCCTTGGTGGTTACATGGC 358
1082 ccagcatctgtctgttgacagcaggtaccagcagtgatgagaggtttgggacctga 1141
359  CCCAGCATCTGTCTGTGGACAGCAGGTACCAGCAGTGGATGGAGAGGTTTGGGCTGA 418
1142 caccagcacttgctctgaatgagaactgtgctcagttcacaaacttcgagcacaa 1201
419  CACCCAGCAGCTTGGTCTGAATGAGAACTGTGCTCAGTTTCAACAGCTTGGCAGCCAAA 478
1202 gattcaaacccagctcaactcatccaccgagacatcttcccctctcaccagtttcg 1261
479  GATTCACAAACCCAGCTCAACTCACTCAACCCGGACATCTTCCCCCTGCTCACCAGTTTCCG 538
1262 ctgtaagaaggaggccccaccctcagtgcccatggttcagggtgaatgcctctcaa 1321
539  CTGTAAGAAGGAGGAGGCCACCCCTCAGTGTGCCATGTTTCAGGTTGAATGCTCCTCAA 598
1322 gtaccagctccgcc 1336
599  GTACCAGCTCCGTC 613

RESULT 28
LOCUS      BE883616          370 bp      mRNA          EST          27-SEP-2000
DEFINITION 601508091F1 NIH_MGC_71 Homo sapiens cDNA clone IMAGE:3909527 5',
            mRNA sequence.
ACCESSION  BE883616
VERSION    BE883616.1 GI:10332392
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 370)
AUTHORS   NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE     National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL   Unpublished (1999)
COMMENT   Contact: Robert Strausberg, Ph.D.
            Tel: (301) 496-1550
            Email: Robert.Strausberg@nih.gov
            Tissue Procurement: ATCC
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLCW710 row: f column: 24
            High quality sequence stop: 370.
            Location/Qualifiers
                1..370
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="IMAGE:3909527"
                /clone_lib="NIH_MGC_71"
                /tissue_type="leiomyosarcoma"
                /lab_host="DH10B (phage-resistant)"
                /note="Organ: uterus; Vector: pCMV-SPORT6; Site 1: NotI;
                Site 2: SalI; Cloned unidirectionally. Primer: Oligo dT.
                Average insert size 2.1 kb."
                Average insert size 2.1 kb.
                94 a 100 c 106 g 70 t

BASE COUNT 94 a 100 c 106 g 70 t
ORIGIN
Query Match 12.5%; Score 370; DB 137; Length 370;
Best Local Similarity 100.0%; Pred. No. 2.1e-184;
Matches 370; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2032 tggctattccgggacaccatccctgcaggtctgtccggatgggaaagatgcca 2091
|||||
1 TGCTCTATTCGGGGACACCATGCCCTGCGAGCTCTGTGTCGGATGGGAAAGATGCCA 60
|||||
```


Tel: 410 955 4678
Fax: 410 614 0827

Email: jeremy_nathans@mail.bs.jhu.edu
Clones from this library are NOT available.
PCR PRIMERS

FORWARD: CTTTGGACCAAGTTCCAGCTGGTTAAGT

BACKWARD: GAGTGGCTATGATGATTTCTTCCAGGTTAA

Seq primer: GGGTAAAGCAAAAGAAAT.

Location/Qualifiers

FEATURES

source

1. .855
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Human retina cDNA randomly primed sublibrary"
/sex="mixed (males and females)"
/tissue_type="retina"
/dev_stage="adult"
/lab_host="E. coli strain K802"
/note="organ: eye; Vector: lambda gt10; Site_1: EcoRI;
Site_2: EcoRI; The library used for sequencing was a
sublibrary derived from a human retina cDNA library.
Inserts from retina cDNA library DNA were isolated,
randomly primed, PCR amplified, size-selected, and cloned
into lambda gt10. Individual plaques were arrayed and
used as templates for PCR amplification, and these PCR
products were used for sequencing."

BASE COUNT 130 a 142 c 162 g 150 t 271 others

ORIGIN

Query Match 11.9%; Score 351; DB 146; Length 855;
Best Local Similarity 100.0%; Pred. No. 2.6e-174; Mismatches 0; Indels 0; Gaps 0;
Matches 351; Conservative 0

995 cttcattcaaccatctgtgagaatccaccttccagaggtaccaggaaggcagatgc 1054
429 CTTCAATCAACCCATCTGTGAGAAATGCCACCTTTCAGAGGTACCAAGGAAGCAGATGC 370
1055 ccccggtgacctgtgtgttcacatgccccagcatctgtgtgtgacagcaggtacca 1114
369 CCCCCTGGGCTTGGTGGTTCACATGCCCCCAGCATCTGTGCTTGTGGACAGCAGGTACCA 310
1115 gcagtgatggagaggtttgggctgacacccagcactgtgtcgaatgagaaactgtgc 1174
309 GCAGTGGATGGAGAGGTTGGGCTGCACACCCAGCAGCTGGTCTGAATGAGAACTGTGC 250
1175 ctgagttcaaaccttcgagccacagattcaaacccagctcaacctcatccaccgga 1234
249 CTCAGTTCACAACTTCGCAGCCACCAAGATTCAACCCAGCTCAACCTCATCCACCCGGA 190
1235 catctccccctctcacacagtttcgctgtgaagaaggaggccccaccctcagttgccc 1294
189 CACTTCCCTCTCTCACCAGTTTCGCTGTGAAGAAGAGGGCCCCCACCCTCAGTGTGCC 130
1295 catggttcaggggtgaatgctctcagtcacagctccgtcccgaggaggga 1345
129 CATGGTTCAGGGTGAATGCTCTCTCAAGTACCAGCTCCGTCCTCCAGGAGGGA 79

RESULT 32

LOCUS BE892893 790 bp mRNA EST 29-SEP-2000
DEFINITION 601435738F1 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:3920792 5',
mRNA sequence.

ACCESSION BE892893

VERSION BE892893.1 GI:10353525

KEYWORDS EST

SOURCE human

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 790)

AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL

COMMENT

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert_Strausberg@nih.gov

Tissue Procurement: ATCC/DCTD/DTF

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov

Plate: LLCW739 row: 1 column: 09

High quality sequence stop: 662.

Location/Qualifiers

FEATURES

source

1. .790
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="IMAGE:3920792"
/clone_lib="NIH_MGC_72"
/tissue_type="melanotic melanoma"
/lab_host="DH10B (phage-resistant)"
/note="organ: skin; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally. Primer: Oligo dI.
Average insert size 2 kb. Library constructed by Life
Technologies."

BASE COUNT 185 a 221 c 220 g 164 t

ORIGIN

Query Match 11.7%; Score 347; DB 137; Length 790;
Best Local Similarity 100.0%; Pred. No. 3.4e-172; Mismatches 0; Indels 0; Gaps 0;
Matches 347; Conservative 0

953 tcttggtgctgtttgtgtgtgtgtagaatgtccagatgaagcttcaatccaccatctg 1012
Db 1 TCTTGTTGCTGCTTTTGTGTGTGTAATGTCCAGATGAAGCTTCATTCAACCCATCTG 60
1013 tgagaatgccaccttcagagtgacaaaggaagcagatccccgtggccttggtgtg 1072
Db 61 TGAGAAATGCCACCTTTCAGAGGTACCAAGGAAGCAGATGCCCGTGGCCTTGGTGTG 120
1073 tcacatggccccagcatctgtgctgtggacagcagggtaccagcagtgagagaggtt 1132
Db 121 TCACATGGCCCCAGCATCTGCTGTGTGGACAGCAGGTACAGCAGTGGATGGAGAGTT 180
1133 tgggcttgacacccagcactgtgtgtcctgaatgagaaactgtgctcagttcacacacttg 1192
Db 181 TGGGCTGTACACCCAGCAGCTTGTGCTCTGAATGAGAACTGTGCTCAGTTCAACACCTTCG 240
1193 cagccacaagattcaaacccagctcaacctcaacctcaccacccggacatctccccctgctcac 1252
Db 241 CAGCCACAAGATTCAAAACCCAGCTCAACCTCAATCCACCCGGACATCTTCCCCCTGCTCAC 300
1253 cagttccgctgtaagaagaggggccccaccctcagtggtgcccattgg 1299
Db 301 CAGTTCCGCTGTAGAAGAGGAGGGCCCCACCCTCAGTGTGCCCATGG 347

RESULT 33

LOCUS AA811170/c 396 bp mRNA EST 19-FEB-1998
DEFINITION ob42c03.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1334020 3',
mRNA sequence.

ACCESSION AA811170

VERSION AA811170.1 GI:2880781

KEYWORDS EST

SOURCE human

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 396)

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

JOURNAL COMMENT
 Tumor Gene Index
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 Tel: (301) 496-1550
 Email: Robert_Strausberg@nih.gov
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman, Ph.D., Gerald Marti, M.D.
 CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.
 CDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CCAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bhrp/image/image.html
 Insert length: 885 Std Error: 0.00
 Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 377.
 Location/Qualifiers
 1. .396
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:1334020"
 /clone_lib="NCI-CCAP_GCB1"
 /tissue_type="germinal center B cell"
 /lab_host="DH10B"
 /note="vector: p7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was prepared from human tonsillar cells enriched for germinal center B cells by flow sorting (CD20+, Igd-), provided by Dr. Louis M. Staudt (NCI), Dr. David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was primed with a Not I - oligo(dT) primer
 [5'-TGTTACCAATCTGAATGGAGCGCGCTCATTTTTTTTTTTTTTTT-3']
). Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p7T3 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."
 81 a 102 c 97 g 116 t

BASE COUNT
 ORIGIN

Query Match 11.6%; Score 343; DB 12; Length 396;
 Best Local Similarity 100.0%; Pred. No. 4.2e-170; Indels 0; Gaps 0;
 Matches 343; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2616 ctgaagagcaggtctcccccaggagcagctcagatagtggtatggagctgtgccgagg 2675
 |||||
 2635 CTGAAGAGCAGGTCCTCCAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 2696
 |||||
 2676 cttgggtccacataagcactatgctatagatgcctctttaggactgtgctggcacag 2735
 |||||
 2695 CTGGGCTCCACATAAGCAGTCTATAGATGCTCTTAGGACTGTGTGCTGGCACAG 2736
 |||||
 2736 ccgaggcagagagctccacacggaagcagcagatgaacttaattcattcaagca 2795
 |||||
 2735 CCGGGGCGCAGAGGCTCCACACGGAAGAGAGAGAGAGAGAGAGAGAGAGAGAG 176
 |||||
 2796 gttttaagaagtcttggaacagacgagcagccttctcttaataccagcaagtat 2855
 |||||
 175 GTTTTAAAGAAGCTTTGGAACAGACGCGGACCTTCTCTTAATCCAGCAAGTAT 116
 |||||
 2856 tccctgcacacagacagacagagtaaacaggtatcagtgggtctaatgtccagactt 2915
 |||||
 115 TCCCTGCACACAGAGACAGAGAGTAAACAGGATCAGTGGGTCTAAGTGTCCGAGACT 56
 |||||
 2916 aacgaaaaatatttcagctgcaataaagattagttgcaa 2958
 |||||
 55 AACGAAAAATATTTCAGCTGCAATAAAGATTGAGTTTGCAA 13
 |||||

RESULT 34
 AA634909

LOCUS AA634909 446 bp mRNA EST 21-OCT-1997
DEFINITION ab27H02.r1 Stratagene lung (#937210) Homo sapiens cDNA clone IMAGE:842067 5' similar to SW:YK59-YEAST P36159 HYPOTHETICAL 96.8 KD PROTEIN IN SIS2-MTD1 INTERGENIC REGION. ; mRNA sequence.
ACCESSION AA634909
VERSION AA634909.1 GI:25598123
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 446)
AUTHORS Hillier.L., Allien.M., Bowles.L., Dubuque.T., Geisel.G., Jost.S., Krizman.D., Kucaba.T., Lacy.M., Le.N., Lennon.G., Marra.M., Martin.J., Moore.B., Schellenberg.K., Steptoe.M., Tan.F., Theising.B., White.Y., Wylie.T., Waterston.R. and Wilson.R.
TITLE WashU-NCI human EST Project
JOURNAL Unpublished (1997)
COMMENT Contact: Wilson RK.
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1870
 Fax: 314 286 1870
 Email: estevaton.wustl.edu
 This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
 Seq primer: -28ml3 revl ET from Amersham
 High quality sequence stop: 430.
 Location/Qualifiers
 1. .446
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:842067"
 /clone_lib="Stratagene lung (#937210)"
 /sex="male"
 /dev_stage="72 years"
 /lab_host="SOLR cells (kanamycin resistant)"
 /note="Organ: lung; Vector: pBluescript SK-; Site_1: EcoRI ; Site_2: XhoI; Cloned unidirectionally. Primer: Oligo dt. normal lung. Average insert size: 1.0 kb; Uni-ZAP XR vector; -5' adaptor sequence: 5' GAATTCGGCAGAG 3' -3' adaptor sequence: 5' CTCGATTTTTTTTTTTTTTTT 3'"
 105 a 112 c 131 g 97 t 1 others

BASE COUNT
 ORIGIN

Query Match 11.6%; Score 342; DB 10; Length 446;
 Best Local Similarity 100.0%; Pred. No. 1.4e-169; Indels 0; Gaps 0;
 Matches 342; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1944 tgtatttgaagaggtttcagacctgtctgtgctgctgctgctgctgctgctgctgct 2003
 |||||
 59 TGTGATTGGAAGAGATTTCAGACCTGTGTGTGGCGGACTGCAAGCATGCGTTTGGCTGT 118
 |||||
 2004 gcctgtgtgcacacctgtgctgctgctgctgctgctgctgctgctgctgctgctgct 2063
 |||||
 119 GCGTGTGTGCACACCTGTGTGTGGAAGTGTCTATTTCGGGGGACACATCCCTGGCAG 178
 |||||
 2064 gctctgtgtgcagtgctgctgctgctgctgctgctgctgctgctgctgctgctgctgct 2123
 |||||
 179 GCTCTGTGCGGATGGGGAAGATGCCACCTCTCTGTATACATGAAGCCACCTCGAAGAT 238
 |||||
 2124 ggtttgaagaggaagcagctggaagacacacacacacacacacacacacacacacac 2183
 |||||
 239 GGTGTTGAAGAGGAGCAGTGTGGAAGAGACACACAGCAGCAGTCCCAAGCCATCAGCGTG 298
 |||||
 2184 gggatgcggatgaacgcggaggttcattatctgtaaccattcagcagcgctatgcaag 2243
 |||||
 299 GGGATGCGGATGAACGCGGAGTTCATTATGCTGAACCACTTCAGCAGCGCTATGCCAAG 358
 |||||
 2244 gtccctctcttcagcccccaacttcagcgagaaagtggagtt 2285
 |||||
 359 GTCCCCCTCTTCAGCCCCCACTTCAGCGAGAAAGTGGAGTT 400
 |||||

```
RESULT 35
LOCUS AA679618 452 bp mRNA EST 02-DEC-1997
DEFINITION ag72c12.s1 Gessler Wilms tumor Homo sapiens cDNA clone
IMAGE:1128502 3' similar to SW:YK59_YEAST P36159 HYPOTHETICAL 96.8
KD PROTEIN IN S1S2-MTD1 INTERGENIC REGION. ; mRNA sequence.
ACCESSION AA679618
VERSION AA679618.1 GI:2660140
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 452)
AUTHORS Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin,
J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Rheising, B.,
White, X., Wyllie, T., Waterston, R. and Wilson, R.
WashU-NCI human EST Project
Unpublished (1997)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Possible reversed clone: similarity on wrong strand
Possible reversed clone: polyT not found
Seq primer: -40m13 fwd. Et from Amersham.
FEATURES
source
1..452
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1128502"
/clone_lib="Gessler Wilms tumor"
/sex="pooled (6)"
/lab_host="DH10B"
/note="Vector: pSPORT1; Site_1: SalI; Site_2: NotI; RNA
was prepared from a pool of 6 anonymous Wilms' tumor RNAs.
RNA was prepared by acid-phenol, followed by one round of
oligo dT selection. cDNA library preparation was with
the BRL/life tech. Superscript Plasmid system. An
oligo-dT NotI primer for first strand synthesis generated
gagggcgccc(t)n at the 3' end of the clones. A 5' SalI
adaptor was used with sequence 5'-gtcgaccacgcgtccg-3'.
Resulting cDNAs were size selected (average size 2 kb).
NotI digested, and ligated into NotI/SalI-cut pSPORT1.
Library was constructed by Dr. Manfred Gessler."
BASE COUNT 111 a 120 c 126 g 95 t
ORIGIN
Query Match 11.6%; Score 342; DB 10; Length 452; --
Best Local Similarity 100.0%; Pred. No. 14e-169; Indels 0; Gaps 0;
Matches 342; Conservative 0; Mismatches 0;
251 ccgggactcggcgccgctctacgtctctccgagttcaaccggtatctctcaactg 310
3 111 CCGGGACTCGGGCGCGCGCTCTACGTCCTCCGAGTTCACCGGTATCTTCACTG 170
311 tggagagcgcttcagagactcatgcaggagcacaaagttaaagggttcgcgcctggacaa 370
3 171 TGGAGAAGCGTTTCAGAGACTCATGCAGGAGCACAAAGTTAAAGGTTGCTCGCGTGACAA 230
371 catattcctgacacaatacactggtctaatgttggggcttaagtgggaattcttacc 430
3 231 CATATTCTTCACACGAATGCATGGTCTTAATGTTGGGGGCTTAAGTGAATGATTTAC 290
431 tttaagggaacggggtctccaaaagtgtgtactttctggacctccacaactggaaaata 490
```

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Db 291 TTAAAGGAACCGGCGCTTCCAAAGTGTGTACTTCTGGACCTCCCAACTGGAATA 350
Qy 491 cctcgaagaatcaaaatattttctgttcattgaaagaatagaaactggtgtgagcc 550
Db 351 CCTCGAAGCAATCAAAATATTTCCTGTCATTAAGGAATAGAACTGGCTGTGCGGCC 410
Qy 551 coactctcctccagaaatcagaggatgaacacatgacagtta 592
Db 411 CCACCTCTGCCCCAGAATACGAGGATGAACCAACCATGACAGTTTA 452
RESULT 36
LOCUS BE795434 698 bp mRNA EST 20-SEP-2000
DEFINITION 601592991F1 NIH_MGC_7 Homo sapiens cDNA clone IMAGE:3946774 5',
mRNA sequence.
ACCESSION BE795434
VERSION BE795434.1 GI:10216632
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 698)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: DCTD/BTP
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LLCM807 row: f column: 23
High quality sequence stop: 389.
Location/Qualifiers
1..698
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3946774"
/clone_lib="NIH_MGC_7"
/tissue_type="small cell carcinoma"
/lab_host="DH10B (phage-resistant)"
/note="Organ: lung; Vector: pOTB7; Site_1: XhoI; Site_2:
EcoRI; cDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
```

```
BASE COUNT 201 a 212 c 175 g 110 t
ORIGIN
Query Match 11.5%; Score 341; DB 135; Length 698;
Best Local Similarity 99.7%; Pred. No. 5e-169;
Matches 391; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1529 gatgaagattcgaatgtcagtgccacactgtcaacataagcccgacacgtctctgct 1588
Db 1 GATGAAGATTTCGAATGTCACTGCCACACTTGTCAACAATAAGCCCGACACGCTCTCTGCI 50
Qy 1589 actgactgtggtgagggcacatttggcagctgtgccgtcattacgagaccaggtgga 1648
Db 61 ACTGACTGTGTTGAGGACACGTTTGGGACGCTGTGCCGTCTATTACGAGACACAGTGA 120
Qy 1649 cagggtcctgggacccctggctgtgtgttgtgtccacctgcacgcagatcaccac 1708
```

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|||||
>b 121 CAGGCTCTGGCACCCTGGCTGCTGTTTGTGTCCACCTGCAGCGAGATCACCAC 180
>y 1709 gggcttgcgaagtattctgtcgcagagaaacgccttggcatcttgggaaagccgct 1768
|||||
>b 181 GGGCTTGCACAGTATCTGTCTCAGAGAGAACGCGCTTGGCATCTTTGGGAAAGCCGCT 240
>y 1769 taacccttctgtgtgttgcgcacacaccagctcaaacgctcgaagcctgcagcagtaacaaa 1828
>b 241 TCACCTCTTGTGTGTGTGTCGCCCCCAACAGCTCAAGCGCTCGCTCCAGCAGTACCACAA 300
>y 1829 ccagtgccagagaggtcctgcaccacacatcagatgattcctcgaatgccttcaggaaag 1888
|||||
>b 301 CCAGTGCCAGGAGGTCCTGCACCATCATCATGATGATTCCTGTCGAATGCTTTCAGGAAG 360
>y 1889 ggcctgagatctccagctcctgcagtggaagat 1920
|||||
>b 361 GGCTGAGATCTCCAGTCTCGCAGTGGAAGAT 392

RESULT 37
LOCUS AW407520 345 bp mRNA EST 16-FEB-2000
DEFINITION UI-HF-BM0-adj-b-07-0-UI.r1 NIH_MGC_38 Homo sapiens cDNA clone
IMAGE:3061957 5', mRNA sequence.
ACCESSION AW407520
VERSION AW407520.1 GI:6926577
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 345)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Eco RI site shown at the beginning of the sequence.
Tissue Procurement: Louis M. Staudt, M.D., Ph.D.
cDNA Library Preparation: M.B. Soares Lab
cDNA Library Arrayed by: M.B. Soares Lab
DNA Sequencing by: M.B. Soares Lab
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Seq primer: M13 Forward.
FEATURES
Location/Qualifiers
1..345
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3061957"
/clone_lib="NIH_MGC_38"
/tissue_type="lymph"
/cell_type="germinal center B cells"
/lab_host="DH10B (UTR)"
/notes="Vector: pRT73-Pac; Site_1: NotI; Site_2: Eco RI;
Constructed from size fractionated cytoplasmic mRNA
(2.5-3.5kb). Directionally cloned. Cells provided by Louis
M. Staudt, Ph.D. Library preparation by Maria de Fatima
Bonaldo, Ph.D. and M. Bente Soares, Ph.D."
BASE COUNT 73 a 94 c 82 g 96 t
ORIGIN
Query Match 11.4%; Score 338; DB 90; Length 345;
Best Local Similarity 100.0%; Pred. No. 1.9e-167; Mismatches 0; Indels 0; Gaps 0;
Matches 338; Conservative 0;

>y 2619 aagagcacggtcccccagagcagctcagataggtgtagtgagctgctgcagagctt 2678
|||||
```

```
Db 345 AAGAGCACGGTCCCCAGGAGGACGCTCAGGATAGTGTGATGGAGCTGTGCCGAGGCTT 286
Qy 2679 gggctccacataaagcaactagtctatagatgcctctttaggactggtgcctgcacagcgg 2738
|||||
Db 285 GGGCTCCACACATAAAGCACTAGTCTATAGATGCTCTTAGGACTGCTGCTGCACAGCG 226
Qy 2739 cgggccagagagctccacacggaagcaagcagatgaactaatttcattcctcaagcagtt 2798
|||||
Db 225 CGGGCCAGAGGCTGCCACACGGAAGCAAGCAGATGAACATAATTCATTCAAGCGAGTT 166
Qy 2799 tttaagaagtcttggaaacagacgcgcaccccttccctcctaatccagcaaatgattcc 2858
|||||
Db 165 TTTAAAGAAGTCTTGAAACAGACGCGGCACCTTCTCTTAATCCAGCAAGTATATCC 106
Qy 2859 ctgcacacagagacaagcagagtaaacagagatcaatggctcgaagtcctgcagagactaac 2918
|||||
Db 105 CTGCACACAGAGACAGCAGAGTAAACAGGATCAGTGGTCTTAAGTGTCCGAGACTTAAC 46
Qy 2919 gaaaatagttatttcagctgcaataaagattgattgc 2956
|||||
Db 45 GAAATAGTATTTCAGCTGCAATTAAGATTGAGTTGC 8

RESULT 38
LOCUS BE742908 677 bp mRNA EST 15-SEP-2000
DEFINITION 601574609F1 NIH_MGC_9 Homo sapiens cDNA clone IMAGE:3835658 5',
mRNA sequence.
ACCESSION BE742908
VERSION BE742908.1 GI:10156900
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 677)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Tissue Procurement: DCTD/DRP
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LICM518 row: a column: 03
High quality sequence stop: 672.
FEATURES
Location/Qualifiers
1..677
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3835658"
/clone_lib="NIH_MGC_9"
/tissue_type="adenocarcinoma cell line"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: ovary; Vector: pOTB7; Site_1: XhoI; Site_2:
EcoRI; cDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5',
adaptor: GGCAGAG(G). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT 160 a 173 c 217 g 127 t
ORIGIN
Query Match 11.3%; Score 334; DB 135; Length 677;
Best Local Similarity 99.8%; Pred. No. 2.6e-165;
Matches 454; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
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```

VERSION      RA632118.1  GI:2555532
KEYWORDS     EST.
SOURCE       human.
ORGANISM     Homo sapiens
REFERENCE    1 (bases 1 to 397)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE        National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
              Tumor Gene Index
JOURNAL      Unpublished (1997)
COMMENT      Contact: Robert Strausberg, Ph.D.
              Tel: (301) 496-1550
              Email: Robert.Strausberg@nih.gov
              Emmert-Buck, M.D., Ph.D.
              CDNA Library Preparation: M. Bento Soares, Ph.D.
              DNA Sequencing by: Washington University Genome Sequencing Center
              Clone distribution: NCI-CGAP clone distribution information can be
              found through the I.M.A.G.E. Consortium/LLNL at:
              www-bio.llnl.gov/bbrp/image/image.html
              Insert Length: 1436 Std Error: 0.00
              Seq primer: -40m13 fwd. ET from Amersham
              High quality sequence stop: 383.
              Location/Qualifiers
FEATURES     source
             1..397
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="IMAGE:113137"
                /clone_lib="NCI-CGAP.Br2"
                /sex="female, pooled"
                /tissue_type="breast"
                /lab_host="DH10B"
                /note="Vector: pT73D-Pac (Pharmacia) with a modified
                polylinker; 1st strand cDNA was prepared from pooled bulk
                breast tumor tissue, and was then primed with a Not I -
                oligo(dT) primer. Double-stranded cDNA was ligated to Eco
                RI adaptors (Pharmacia), digested with Not I and cloned
                into the Not I and Eco RI sites of the modified pT73
                vector. This library is the normalized version of
                NCI-CGAP.Br1.1. Library was constructed by Bento Soares
                and M. Fatima Bonaldo."
BASE COUNT   98 a 100 c 111 g 88 t
ORIGIN
Query Match 10.9%; Score 323; DB 10; Length 397;
Best Local Similarity 100.0%; Pred. No. 1.7e-159;
Matches 323; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 251 ccgggactcgggcgcgcgtctctacgtttctccgagttcaaccggtatctctcaactg 310
D 71 CCGGGACTCGGGCGCGCGCTCTACGCTCTCCGAGTTCAACGGGTATCTTCAACTG 130
Y 311 tggagaaggcgttcagagactcatcgaggacacaaagtttaagttgctcgctggacaa 370
D 131 TGGAGAAGGCCTTCAGAGACTCATCGAGGACACAAAGTTAAGGTTCGCTCGCTGGACAA 190
Y 371 catattctgacacgaatgcactggtctcaatgttgggggcttaagtgaatgattcttac 430
D 191 CATATTCTGCACAGATGCACTGCTAATGTTGGGGGCTTAAGTGAATGATCTTAC 250
Y 431 tttaaagaaaccggggtttccaaagtgtgtactttcttgacctccacaaactggaaaata 490
D 251 TTTAAAGAAACCGGGTTCCAAAGTGTGTACTTCTGGACCTCCACAACCTGGAATAA 310
Y 491 cctcgaagcaatcaaatattttctgtccattgaagggaatacactggtgtgcggcc-550
D 311 CCTCGAGCAATCAAAATATTTTCTGGTCCATTGAAGGAATAGAACTGTGTGCGGCC 370
Y 551 ccactctgccccagaatacagg 573
|||||

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Db 371 CCACTCTGCCCAAGATACGAGG 393
RESULT 43
LOCUS    R55841
DEFINITION
Yg89401.r1 Soares infant brain INIB Homo sapiens cDNA clone
IMAGE:40931 5' similar to SP:YK59_YEAST P36159 HYPOTHETICAL 96.8 KD
PROTEIN IN SIS-MTDL INTERGENIC ;, mRNA sequence.
R55841
ACCESSION R55841.1 GI:825947
VERSION R55841.1
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 482)
AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman,
M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,
Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaskis, E., Waterston,
A., Williamson, A., Wohlmann, P., and Wilson, R.
TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Wilton RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Insert Size: 1777
High quality sequence stops: 387 Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1777 Std Error: 0.00
Seq primer: M13RPI
High quality sequence stop: 387.
Location/Qualifiers
FEATURES     source
             1..482
                /organism="Homo sapiens"
                /db_xref="GDB:413472"
                /db_xref="taxon:9606"
                /clone="IMAGE:40931"
                /clone_lib="Soares infant brain INIB"
                /sex="female"
                /dev_stage="73 days post natal"
                /lab_host="DH10B (ampicillin resistant)"
                /note="Organ: whole brain; Vector: Lfamid BA; Site_1: Not
                I; Site_2: Hind III; 1st strand cDNA was primed with a Not
                I - oligo(dT) primer [5'
                AACTGGAAGAATTCGGCGCGCGAGGAATTTTCTTTTCTTTT 3'];
                double-stranded cDNA was ligated to Hind III adaptors
                (Pharmacia), digested with Not I and directionally cloned
                into the Not I and Hind III sites of the Lfamid BA vector.
                Library went through one round of normalization. Library
                constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT   107 a 139 c 131 g 102 t 3 others
ORIGIN
Query Match 10.9%; Score 323; DB 144; Length 482;
Best Local Similarity 100.0%; Pred. No. 1.7e-159;
Matches 323; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1430 gcaggagtcagaggagtgccgaggacgcccagccccgcagagagaaagatcagta 1489
D 43 GCAGGAGTACAGGAGGAGTGCAGGACGCCGCCAGCCAGAGAAAAGAGTCACTA 102
Y 1490 ccagagaatcatcttccttgagacaggtctgccatcccatgaagattcgaaatgcag 1549
D 103 CCCAGAAATCATCTCTCTTGGACACAGGGTCTGCCATCCCATGAAGATTCCAAATGTCAG 162
Y 1550 tggcacactgtcaacataagccccgacacgtctctgctactgactggtggtggggcac 1609
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> 163 TGCCACACTTGTCAACATAAGCCCGACACGTCCTCTGCTACTGACTGTGGTGGGGCAC 222
> 1610 atttgggagctgtgcccattacggagacaggtgacaggggtccctgggacacctggc 1669
> 223 ATTTGGGAGCTGTGCCGTCTATTACGGAGACAGGTGGACAGGGTCCCTGGGCACCTGGC 282
> 1670 tgcgtgtttgtgtccacctgcacgcagatcaccacacggcgttgccaaagtattctgt 1729
> 283 TGTGTGTTTGTGTCCTACCTGTCAGCGAGATCACACACGGGCTTGCCAAAGTATCTTGT 342
> 1730 gcagagagaacgcgcttggcat 1752
> 343 GCAGAGAGACGGCGCTTGGCAT 365

RESULT 44
LOCUS AA233087/c 366 bp mRNA EST 28-FEB-1997
DEFINITION z168g08.r1 Soares_NHMPu_S1 Homo sapiens cDNA clone IMAGE:568606
5', mRNA sequence.
ACCESSION AA233087
VERSION AA233087.1 GI:1856275
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 366)
AUTHORS Hillier,L., Clark,N., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
R., Williamson,A., Woldmann,P. and Wilson,R.
The Washo-Merck EST Project
UNPUBLISHED (1995)
CONTACT: Wilton RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -28m13 rev2 ET from Amersham
High quality sequence stop: 346.
FEATURES
Location/Qualifiers
1..366
/organism="Homo sapiens"
/db_xref="GDB:5562573"
/db_xref="taxon:9606"
/clone_lib="IMAGE:568606"
/clone_lib="Soares_NHMPu_S1"
/tissue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/lab_host="DH10B"
/notes="Organ: mixed (see below); Vector: pT7T3D-Pac
(Pharmacia) with a modified polylinker; Site 1: Not I;
Site 2: Eco RI; Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NbW, pregnant uterus
NbHPU, and fetal heart NbHH19W) were mixed, and ss circles
were made in vitro. Following HAP purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of I.M.A.G.E. clones 260232-265223,
340488-345479, and 484488-489479."
BASE COUNT 83 a 90 c 110 g 83 t
GIGIN

Query Match 10.6%; Score 315; DB 4; Length 366;
Best Local Similarity 99.7%; Pred. No. 2.9e-155;
Matches 365; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 931 aagagctgtactctccagatctgctgctgcttctggtgtagaattgctcagatg 990.
DB 366 AAGAGCTGTGTACTTCCACAGATCCTGGTGTCTTTTGTGGTAGAATGTCCAGATG 307
QY 991 aaagcttcattcaacccatctgtgagaatgccaccttccagaggtaccagaaggaagcag 1050
DB 306 AAGCTTCATTCAACCCATCTGTGAGAAATGCCACTTTCAGAGGTACCAAGAAATGCAG 247
QY 1051 atgcccccgctggccttgggttgcacatggccccagacatctgtctgtgacagcaggt 1110
DB 246 ATGCCCGCTGGCTTGGTGTTCACATGGCCCCAGCATCTGTCTTGTGACAGAGGT 187
QY 1111 accagcagtgatgagaggtttgggctgacacccagcacttggctctgaatgagaact 1170
DB 186 ACCAGCAGTGGATGGAGAGGTTTGGGCTTGACACCCAGCACTTGGTCTGTGAATGAACT 127
QY 1171 gtgctcagtttcacaaccttcgcagcagcacaagattcaaacccagctcaacctctccacc 1230
DB 126 GTGCTCAGTTTCAACACTTTCGCAGCCACAAGATTCAAAACCAGCTCAACCTCATCCACC 67
QY 1231 cggacatcttccccctgctcaccagtttccgctgtaagaagaggagggccccacctcagt 1290
DB 66 CGGACATCTTCCCTCTGCTCACCAGTTTCCGCTGTAAAGAGGAGGGCCCCACCTCAGTG 7
QY 1291 tgcaca 1296
DB 6 TGCCCA 1

RESULT 45
LOCUS BE902696 735 bp mRNA EST 29-SEP-2000
DEFINITION 601677393F1 NIH_MGC_21 Homo sapiens cDNA clone IMAGE:3959926 5',
mRNA sequence.
ACCESSION BE902696
VERSION BE902696.1 GI:10393148
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 735)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
NATIONAL INSTITUTES OF HEALTH, MAMMALIAN GENE COLLECTION (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
Plate: LLCM841 row: j column: 23
High quality sequence stop: 732.
FEATURES
Location/Qualifiers
1..735
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="IMAGE:3959926"
/clone_lib="NIH_MGC_21"
/tissue_type="choriocarcinoma"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: placenta; Vector: pOTB7; Site 1: XhoI;
Site 2: EcoRI; cDNA made by oligo-dt priming.
Directionally cloned into EcoRI/XhoI sites using the
following 5' adaptor: GGCAGCAG(G). Size-selected by
for average insert size 1.8kb. Library constructed by
Ling Hong in the laboratory of Gerald M. Rubin (University
of California, Berkeley) using ZAP-CDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT 168 a 206 c 203 g 157 t 1 others

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479 actggaaaaataactcgaagaacaataaataattttcttgcctccattgaaaggaataagaact 538
|||||
339 ACTGGAAAAATACCTCGAAGCAATCAAAATATTTCTGGTCCATTGAAAGGAATAGAACT 398
|||||
539 gctgtgcggccactctcccaagaatacagagatgaaccatacacagttaccagat 598
|||||
399 GGTGTGCGGCCCCACTCTCCCCAGAAATACGAGGATGAACCATGACATTTACAGAT 458
|||||
599 ccccatcacagtgaacagagggggaagcaccacaccatggcagagtcag 651
|||||
459 CCCATACACAGTGAACAGAGGAGGGGAAACACCAACCATGCGAGAGTCCAG 511
|||||

RESULT 49
LOCUS AW994476 297 bp mRNA EST 05-JUN-2000
DEFINITION RC3-BN0036-260400-015-e06 BN0036 Homo sapiens cDNA, mRNA sequence.
ACCESSION AW994476
VERSION AW994476.1 GI:8254699
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 297)
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare
M.J., Soares,F., Brentani,R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.
Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?t1=st2-RC3-BN0036-260
400-015-e06&t3=2000-04-26&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 9
High quality sequence stop: 297.
FEATURES
Location/Qualifiers
1..297
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="BN0036"
/dev_stage="Adult"
/note="Organ: breast_normal; Vector: puc18; Site_1: Smal;
Site_2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."
60 a 93 c 82 g 62 t
BASE COUNT 60 a 93 c 82 g 62 t
ORIGIN
Query Match 9.8%; Score 291; DB 97; Length 297;
Best Local Similarity 100.0%; Pred. No. 1.5e-142;
Matches 291; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
1567 taagcccccacagctctcgtactgactgctggtgagggcacatttggcagctgtgcc 1626
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Db 7 TAAGCCCCACACGCTCTGCTACTGACTGCTGGTGGAGGCACATTGGGCACGCTGGCC 55
QY 1627 gtattacggagaccaggtggacagaggtctggcaccctggctgtgtgttgttctccc 1686
|||||
Db 67 GTCATTACGGAGACCAAGTGGACAGGGTCTCTGGCACCCCTGGCTGTGTTGTGTCCC 126
QY 1687 acctgacacagatcacacacaggggtttcccaagtattctgtgcagagagaacgcct 1746
|||||
Db 127 ACCTGCACGACATCACACACGGGTTGCCAAGTATCTTGCTGCAGAGAGAACGGCCT 186
QY 1747 tggcatctttgggaagcgcttcaccccttgcgtggttgcgttgcgttgcgttgcgttgc 1806
|||||
Db 187 TGGCATCTTTGGGAAGCGCTTCACCCCTTGTGCTGGTGGCCCCCACCACGCTCAAG 246
QY 1807 cctggtccagcagtcaccacacacagtcgacagaggtcctgcaccacatca 1857
Db 247 CCTGGTCCAGCAGTACCACACACAGTGCAGGAGTCTCTGCACCATCA 297

RESULT 50
LOCUS AII174501 404 bp mRNA EST 07-OCT-1998
DEFINITION an42e05.s1 Gessler Wilms tumor Homo sapiens CDNA clone
IMAGE:1701344 3' similar to SW:YK59_YEAST P36159 HYPOTHETICAL 96.8
KD PROTEIN IN S152-MTD1 INTERGENIC REGION. ; mRNA sequence.
ACCESSION AII174501
VERSION AII174501.1 GI:3721354
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 404)
Hillier,L., Allen,M., Bowles,L., Duboue,T., Geisels,G., Jost,S.,
Krisman,D., Kucaba,T., Lacy,M., Le,N., Lennon,G., Marra,M., Martin
J., Moore,B., Schellenberg,K., Steptoe,M., Tan,F., Theising,B.,
White,Y., Wyllie,T., Waterston,R. and Wilson,R.
WashU-NCI human EST Project
Unpublished (1997)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LML; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Possible reversed clone: similarity on wrong strand
Possible reversed clone: polyt not found
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 391.
FEATURES
Location/Qualifiers
1..404
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="IMAGE:1701344"
/sex="pooled (6)"
/lab_host="DH10B"
/note="Vector: pSPORT1; Site_1: SalI; Site_2: NotI; RNA
was prepared from a pool of 6 anonymous Wilms' tumor RNAs.
RNA was prepared by acid-phenol, followed by one round of
oligo dr selection. CDNA library preparation was with
the BRL/Life Tech. Superscript Plasmid system. An
oligo-dr NotI primer for first strand synthesis generated
gcggcgcccc(t)n at the 3' end of the clones. A 5' SalI
adaptor was used with sequence 5'-gtcagccacgcgtccg-3'.
Resulting cDNAs were size selected (average size 2 kb),
NotI digested, and ligated into NotI/SalI-cut pSPORT1.
Library was constructed by Dr. Manfred Gessler."
97 a 104 c 115 g 87 t
BASE COUNT 97 a 104 c 115 g 87 t
ORIGIN

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SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 397)
AUTHORS Hillier, L., Clark, N., Dubucque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J., Rifkin, L., Rohlfs, T., Soares, M., Tan, F., Trevaskis, E., Waterston, R., Williamson, A., Wohlmann, P. and Wilson, R.
TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Wilton RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
Possible reversed clone: similarity on wrong strand
Possible reversed clone: polyT not found
Insert Length: 678 Std Error: 0.00
Seq primer: mob.REGA+ET.
FEATURES
source
Location/Qualifiers
1..397
/organism="Homo sapiens"
/db_xref="GDB:1259575"
/db_xref="taxon:9606"
/clone="IMAGE:32191"
/clone_lib="Soares.parathyroid_tumor_NBHPA"
/tissue_type="parathyroid tumor"
/dev_stage="adult"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: parathyroid gland; Vector: pTTT3D (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer
[5'-TGTTACCAATCTGAAGTGGGCGCGCCACCAATTTTTTTTTTTTTTTT
TTTTT-3'] double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pTTT3 vector (Pharmacia). Library went through one round of normalization to a Cot = 5. Library constructed by Bento Soares and M. Fatima Bonaldo. RNA from sporadic parathyroid adenomas was kindly provided by Dr. Stephen Marx, National Institute of Diabetes and Digestive and Kidney Diseases, NIH."
BASE COUNT 97 a 101 c 112 g 83 t 4 others
ORIGIN
Query Match 9.6%; Score 285; DB 146; Length 397;
Best Local Similarity 100.0%; Pred. No. 2.3e-139;
Matches 285; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
252 cggagctcggcgccgctctacgtcttctcgaagttcaacgggtatcttcaactgt 311
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Db 352 CTCGAAGCAATCAAAATATTTTCTGTCATTGAAAGGAATAGAA 396
RESULT 55
AW248468/c
LOCUS
DEFINITION 2820640.3prime NIH_MGC_7 Homo sapiens cDNA clone IMAGE:2820640 3',
mrna sequence.
ACCESSION AW248468
VERSION AW248468.1 GI:5591461
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 394)
AUTHORS NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Other ESTs: 2820640.5prime
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: DCTD/DTP CDNA Library Preparation: Ling Hong/Rubin Laboratory CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Berkeley MGC sequencing project
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www.bio.llnl.gov/bbrp/image.html Base Calling / Quality Scores: PHRED from University of Washington Genome Center Trimming: cross_match from University of Washington Genome Center PHRAP suite. Poly-T Identification: patmatch.pl from Berkeley Drosophila Genome Project. University of Washington Genome Center: http://www.genome.washington.edu Polyadenylation: Based upon the presence of a XhoI site followed by a run of 14 or more T residues at the beginning of the sequence, this cDNA insert was polyadenylated.
Plate: L10M4 row: L column: 17
High quality sequence stop: 213.
Location/Qualifiers
1..394
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2820640"
/clone_lib="NIH_MGC_7"
/tissue_type="small cell carcinoma"
/cell_line="MGC3"
/lab_host="DH10B (phage-resistant)"
/note="Organ: lung; Vector: pOTB7; Site_1: XhoI; Site_2: EcoRI; CDNA made by oligo-dT priming Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."
BASE COUNT 80 a 103 c 97 g 114 t
ORIGIN
Query Match 9.6%; Score 284; DB 88; Length 394;
Best Local Similarity 99.5%; Pred. No. 7.7e-139;
Matches 384; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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274 GCATGCTATAGATGCTCTTAGACTGGTGGCTGGACACCGCGGGCCAGGAGTT 215
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2753 gccacaggaagcaagcagatgaactaatattcatttcaagcaggttttaagaagtctt 2812
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2813 ggaacacagcggcgacacctttctctctaatccagcaaaagtattccctgcacaccagaga 2872
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2933 agtgcgaataaagattgattgtgcaa 2958
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34 AGCTGCAATAAAGATTGAGTTGCAA 9
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RESULT 56
LOCUS AI803400/c
DEFINITION tc42f03.x1 Soares_total_fetus_Nb2HF8_9w Homo sapiens cDNA clone
IMAGE:2067293 3', mRNA sequence.
ACCESSION AI803400
VERSION AI803400.1 GI:5368962
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 489)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1275 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 446.
Location/Qualifiers
1..489
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Soares_total_fetus_Nb2HF8_9w"
/dev_stage="8-9 weeks"
/lab_host="DH10B"
/notes="Vector: pT73D-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was prepared from mRNA obtained from pooled 8-9 week
(total) fetus material with a Not I - oligo(dT) primer [5'
TGTTACATCTCACTGAGTGGAGCGCGCTTAATTTTATTTTATTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT 94 a 134 c 124 g 137 t
ORIGIN

Query Match 9.6%; Score 283; DB 25; Length 489;
Best Local Similarity 99.2%; Pred. No. 2.6e-138;
Matches 483; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

2472 cctcagcagaagcggccccacagagagccacagagcgaaggtcagagcccgagta 2531
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Db 489 CCTACGACAGAAGCGGGCCACACACAGAGGAGCCACAGGGCCAAAGGTCAGAGCCCAAGTGA 430
QY 2532 agatctggagagaccctgaactcagaaggtgtgtgtcttcttcccccacgacagccctt 2591
Db 429 AGATCTGGGAGAGCCCTGAACCTCAGAAGGCTGTGTGCTTCTGCCCCACACGACCCCGT 370
QY 2592 atctgcctccctctgctggttagaagctgaagcagcagcaggttccccccagagcagctcaggt 2851
Db 369 AYTGGCCCTCTCTGCTGGTGAAGCTGAAGACGACGCTCCCGGAGGAGCGAGCTCAGGAT 310
QY 2652 aggtggtatggagctgtgcccagaggttgggctccacataagcaactagtcttagatgcc 2711
Db 309 AGGTGATATGGAGCTGTGCCGAGGCTTGGGTGCCACATAAAGCACTAGTCTATAGATGCC 250
QY 2712 tcttagaagctgtgctgcacagcgcgcccagagaggtccacacggaagcaagcag 2771
Db 249 TCTTAGGACTGTGCTGCTGGCAGAGCTGCGGGCCAGAGGCTGCCACACGAGCAAGCAG 190
QY 2772 atgaactaatcttcatttcaggcagtttttaagaagttttgaaacagacgagcgccacc 2831
Db 189 ATGACTAATTTCAATTCAGGCAGTTTTTAAGAGAGTCATGGAAACACACGCGCGCACC 130
QY 2832 ttctcttaacacagcaagtgattccctgcacacagacagacagcagagtgtaacaggtc 2891
Db 129 TTTCCCTCTAATCAGCAAAATGATTCCTCCACACAGAGACAAGCAGAGTAGTAACAGGATC 70
QY 2892 agtgggttaagtgctccgagacttaacgaaataagtatttcagctgcaataaagattgag 2951
Db 69 AGTGGGTCTAAGTGTCGAGACTTAACGAAATAGTATTTCAGCTCAATTAAGATTGAG 10
QY 2952 ttgtgcaa 2958
Db 9 TTGCA 3

RESULT 57
LOCUS T34216
DEFINITION EST64346 Human White blood cells Homo sapiens cDNA 5' end similar
to None, mRNA sequence.
ACCESSION T34216
VERSION T34216.1 GI:616314
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 307)
EST64346 Human White blood cells Homo sapiens cDNA 5' end similar
to None, mRNA sequence.
ACCESSION T34216
VERSION T34216.1 GI:616314
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 307)
ADAMS, M.D., Kerlavage, A.R., Fleischmann, R.D., Fuldner, R.A., Bult
, C.J., Lee, N., Kirkness, E.F., Weinstock, K.G., Gocayne, J.D., White
, O., Sutton, G., Blake, J.A., Brandon, R.C., Chiu, M.-W., Clayton, R.A.,
Cline, R.T., Cotton, M.D., Earle-Hughes, J., Fine, L.D., FitzGerald
, L.M., FitzHugh, W.M., Fritchman, J.L., Geoghagen, N.S.M., Glodek, A.,
Gnehm, C.L., Hanna, M.C., Hedblom, E., Hinkley, P.S., Kelley, J.M.,
Klimek, K.M., Kelley, J.F., Liu, L.-I., Marmaros, S.M., Merrick, J.M.,
Moreno-Palauques, R.F., McDonald, L.A., Nguyen, D.T., Pellegrino, S.M.,
Phillips, C.A., Ryder, S.E., Scott, J.L., Saudke, D.M., Shirley, R.,
Small, K.V., Spriggs, T.A., Utterback, T.R., Weidman, J.F., Li, Y.,
Bednarek, D.P., Cao, L., Cepeda, M.A., Coleman, T.A., Collins, E.-J.,
Dimke, D., Feng, P., Ferrie, A., Fischer, C., Hastings, G.A., He, W.-W.,
Hu, J.-S., Greene, J.M., Gruber, J., Hudson, P., Kim, A., Kozak, D.L.,
Kunsch, C., Li, H., Li, H., Meissner, P.S., Olsen, H., Raymond, L., Wei
, Y.-F., Wing, J., Xu, C., Yu, G.-L., Ruben, S.M., Dillon, P.J., Fannon
, M.R., Rosen, C.A., Haseltine, W.A., Field, S.C., Fraser, C.M. and
Venter, J.C.
Initial Assessment of Human Gene Diversity and Expression Patterns
Based Upon 83*Million Basepairs of cDNA Sequence
Nature 377, 3-174 (1995)
JOURNAL 96026280
MEDLINE
COMMENT Other ESTs: THC15444
Contact: Venter, JC
932 Clopper Rd, Gaithersburg, MD 20878
```


Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 278)
Dias Neto,E., Garcia Correa,R., Vexijovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zaço,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Masukuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare ,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.
Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
20202663
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(<http://www.ludwig.org.br/scripts/gethtml2.pl?tl-st2=IL0-ST0002-160>)
599-0034t3-1999-05-16&t4=1)
Seq primer: puc 18 forward
High quality sequence stop: 278.

[illegible]

Accession	Gene	Length (bp)	Source
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U00085	Stratagene	461	EST
U00086	Stratagene	461	EST
U00087	Stratagene	461	EST
U00088	Stratagene	461	EST
U00089	Stratagene	461	EST
U00090	Stratagene	461	EST

IMAGE:844019 5', mRNA sequence.
AA635046
VERSION AA635046.1 GI:2558260
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 461)
Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisels, G., Jost, S.,
Krisman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Mar
J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B.,
White, Y., Wylie, T., Waterston, R. and Wilson, R.
WashU-NCI human EST Project
Unpublished (1997)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (infoimage.llnl.gov) for further information.
Insert Length: 694 Std Error: 0.00
Seq primer: -28mi3 rev1 ET from Amersham
High quality sequence stop: 453.

BASE COUNT	110 a	120 c	159 g	72 t	
ORIGIN					
Query Match	9.3%;	Score 276;	DB 10;	Length 461;	
Best Local Similarity	99.7%;	Pred. No. 1.3e-134;			
Matches 326;	Conservative	0;	Mismatches 1;	Indels	0; Gaps
Qy	2410	tgcggcaggtgcgggcggccctctctgtccaggagctggcaggcgccctggagagatgggg	2469		
Db	42	TGCGGCAGGTGCGGGCGGCCCTCTGTCTCCAGGAGCTGGCAGGGCGGCTGGAGGATGGG	101		
Qy	2470	agcctcagcagaagcgggcccacacagagagccacaggccaaagaaggtcagagcccagt	2529		
Db	102	AGCCTCAGCAGAAAGCGGGCCCAACACAGAGAGGCCACAGCCGCAAGAGGTGACAGCCCCAGT	161		
Qy	2530	gaagatctgggagacccctgaactcgaagaggtgtgtgtctcttctgccccacgcacgaccc	2589		
Db	162	GAAGATCTGGGAGACCCCTGAACTCAGAAGGGTGTGTGTCTTCTGCCCCACGCACGCACCC	221		
Qy	2590	gtatctgccctctctgtgtagaagctgaagacacaggtccccccaggagggcagctcagg	2649		
Db	222	GTATCTGCCCTCTCTGTCTGGTAGAAGCTGAAGAGCACGGTCCCCCAGGAGGCAGCTCAGG	281		
Qy	2650	atagggtgtagagagctgtgcgagggcttgggtccccacataaagcactagtcttatagatg	2709		
Db	282	ATAGGTGGTAGAGCTGTGCCAGGCTTGGCTCCCACTAAGCACTAGTCTATAGATG	341		
Qy	2710	cctcttagcactggtgcctggcacgc	2736		
Db	342	CCCTTTAGGACTGGTGCCTGGCAGCAGC	368		


```

RESULT 61
LOCUS AW175581 364 bp mRNA EST 16-NOV-1999
DEFINITION OVO-BT0041-030999-013-e09 BT0041 Homo sapiens cDNA, mRNA sequence.
ACCESSION AW175581
VERSION AW175581.1 GI:6441618
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 364)
AUTHORS HCSP http://www.ludwig.org.br/ORESTES.
TITLE The FAPESP/LICR Human Cancer Genome Project
JOURNAL Unpublished (1999)
COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?CI=QV0&t2=QV0-BT0041-
030999-013-e09&t3=1999-09-03&t4=1)
Seq primer: puc 18 forward
High quality sequence stop: 75
High quality sequence stop: 364.
FEATURES
Location/Qualifiers
1..364
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="BT0041"
/dev_stage="Adult"
/note="Organ: breast; Vector: puc18; Site_1: SmaI; Site_2:
SmaI; A mini-library was made by cloning products derived
from ORESTES PCR (U.S. Letters Patent application No. 196
716 - Ludwig Institute for Cancer Research) profiles
into the pUC 18 vector. Reverse transcription of tissue
mRNA and cDNA amplification were performed under low
stringency conditions."
BASE COUNT 75 a 93 c 118 g 78 t
ORIGIN
Query Match 9.3%; Score 274; DB 87; Length 364;
Best Local Similarity 100.0%; Pred. No. 1.5e-133;
Matches 274; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1025 ctttcagaggtaccagaagaagcagatgcccccgtggccttggttggttcacatgcccc 1084
13 364 ctttcagaggtaccagaagaagcagatgcccccgtggccttggttggttcacatgcccc 305
17 1085 agcatctgtgttggtgacagcaggtaccagcagtggtgagaggttttgggacctgacac 1144
20 304 agcatctgtgttggtgacagcaggtaccagcagtggtgagaggttttgggacctgacac 245
23 1145 ccagcactgtgttcgagatggaactgtgttcagttcacacaccttcagccacaagat 1204
26 244 ccagcactgtgttcgagatggaactgtgttcagttcacacaccttcagccacaagat 185
29 1205 tcaaacccagctcaacccctcaccgccgacatctcccccctgctcaccagtttccgctg 1264
32 184 tcaaacccagctcaacccctcaccgccgacatctcccccctgctcaccagtttccgctg 125
35 1265 taagaagagggcccccaccctcagtggtgcccattg 1298
38 124 taagaagagggcccccaccctcagtggtgcccattg 91

```

```

RESULT 62
LOCUS AA994126 457 bp mRNA EST 27-AUG-1998
DEFINITION Ou38b06.s1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
IMAGE:1628531 3' similar to SW:YK59.YEAST P36159 HYPOTHETICAL 95.8
KD PROTEIN IN SIS2-MTD1 INTERGENIC REGION. ; mRNA sequence.
ACCESSION AA994126
VERSION AA994126.1 GI:3180671
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 457)
AUTHORS NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.llni.gov) for further information.
Insert Length: 1193 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 439.
FEATURES
Location/Qualifiers
1..457
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/note="Organ: pooled; Vector: pT73D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NBHL19W, testis NHT, and B-cell
NCI-CGAP-GCB1) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 682632-687239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo.
BASE COUNT 115 a 119 c 127 g 96 t
ORIGIN
Query Match 9.2%; Score 273; DB 14; Length 457;
Best Local Similarity 100.0%; Pred. No. 5.2e-133;
Matches 273; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 279 ttctccgagttcaaccggtatctctcaactgtgagaagcggttcagagactcatgcaag 338
Db 138 TTCTCCGAGTTCAACCCGGTAICTCTTCACTGTGAGAGAGCGCTTCAGAGACTCATGAC 197
QY 339 gagcacaagttaaaaggttgctgcctggacaacatatctctgacacgaatgcactggtct 398
Db 198 GAGCAAGTTAAAGGTTGCTCGCTGGACAACATATTCTTGACACGAATGCACCTGGTCT 257
QY 399 aatgttggggccttaagtgggaatgattcttcttaagggaacccggccttccaaagtgt 458
Db 258 AATGTTGGGGCCTTAAGTGGAAATGATTCTTACTTTAAAGGAAACCGGCTTCCAAGTGT 317
QY 459 gtactttctgacctcccaactgcaaaaatactctcgaagcaatcaaaatatttctgt 518
Db 318 GTACTTTCTGACCTCCCAACTGGAAAAAATACCTCGAAGCAATCAAAATATTTCTGTGT 377
QY 519 ccattgaagaagaatagaactgctgtgcggccc 551
Db 378 CCATTGAAGGAATAGAACTGGCTGTGCGGCC 410

```

```
RESULT 63
OCUS BE744876 984 bp mRNA EST 15-SEP-2000
DEFINITION 601576324F1 NIH_MGC_9 Homo sapiens cDNA clone IMAGE:3837533 5',
        mRNA sequence.
ACCESSION BE744876
VERSION BE744876.1 GI:10158868
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 984)
NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/...
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: DCTD/DTP
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at: Image.llnl.gov
Plate: LHC522 row: o column: 06
High quality sequence stop: 726.
Location/Qualifiers
1..984
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3837533"
/clone_lib="NIH MGC 9"
/tissue_type="adenocarcinoma cell line"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: ovary; Vector: pOTB7; Site_1: XhoI; Site_2:
EcoRI; cDNA made by oligo-dT priming. Directionally
cloned into EcoRI/XhoI sites using the following 5'
adaptor: GGACGAG(G). Size-selected >500bp for average
insert size 1.8kb. Library constructed by Ling Hong in
the laboratory of Gerald M. Rubin (University of
California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life technologies)."
```

```
BASE COUNT 227 a 259 c 316 g 182 t
ORIGIN
Query Match 9.0%; Score 266; DB 135; Length 984;
Best Local Similarity 99.5%; Pred. No. 2.8e-129;
Matches 436; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 966 ttgtggtgtagaagtcagatgaagctcattcaacccatctgtgagaatgccacc 1025
DB 244 TTGTGGTGTGATGATGTCAGATGAAGCTTCATCAACCCATCTGTGATGATGCCACC 303
QY 1026 ttccagggtaccagaagaaaggcagatgcccccggtggttgggttcacatggcccca 1085
DB 304 TTTCAGAGGTACCAAGGAAGGAGAGATGCCCGCTGGCTTGGTGTTCATATGGCCCCA 363
QY 1086 gcatctgtcttggagcaacagattaccacagtcagtgatgagaggttggcctgacacc 1145
DB 364 GCATCTGTCTGTGGACACAGATGACAGGAGGAGGAGG-RTGGGCGCTGACACC 422
QY 1146 cagcacttggtcctgaatgagaactgctcagttcacaaccttcgacccacaagatt 1205
DB 423 CAGCACTTGTCTCTGAATGAGAACTGTGCTCTCAGTTCACAACTTCGCACCCACAGATT 482
QY 1206 caaaccagctcaactcatccaccgggacatcttcccctgtctcaccagttccctgtgt 1265
DB 483 CAAACCCAGCTCAACCTCATCCACCCGGACATCTTCCCCCTGTCTCACCAGTTCCGCTGT 542

QY 1266 aagaagagggcccccacccctcagtggtgccatggttcagggtgaatgcctccaagtac 1325
DB 543 AAGAGAGGGGGCCCCACCCCTCAGTGTGCCCATGGTTCAGGGTGAATGGCTCTCAAGTAC 602
QY 1326 cagctccgtcccccagggagtgaggagaggtgacattacttacttgcaatcctcaggaa 1385
DB 603 CAGCTCCGTCCCGAGGAGGAGTGGCAGAGGATGCCATTACTTACTTCAATCCTGAGGAA 652
QY 1386 ttcattagtgaggcgcgtg 1403
DB 663 TTCATAGTTGAGGCGCTG 680

RESULT 64
LOCUS AI033108 410 bp mRNA EST 28-AUG-1998
DEFINITION ow98g08.sl Soares_fetal_liver_spleen_lNFLS_S1 Homo sapiens cDNA
        clone IMAGE:1654910 3', mRNA sequence.
ACCESSION AI033108
VERSION AI033108.1 GI:3254061
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 410)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 773 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 368.
Location/Qualifiers
1..410
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1654910"
/clone_lib="Soares_fetal_liver_spleen_lNFLS_S1"
/sex="male"
/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/notes="Organ: Liver and Spleen; Vector: pT7T3D (Pharmacia)
with a modified polylinker; Site_1: Pac I; Site_2: Eco RI;
This is a subtracted version of the original Soares fetal
liver spleen lNFLS library. 1st strand cDNA was primed
with a Pac I - oligo(dT) primer [5',
AACTGGAAGAATAATAAGATCTTTTTTTTTTTTTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pT7T3 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
```

```
BASE COUNT 88 a 110 c 101 g 111 t
ORIGIN
Query Match 8.6%; Score 255; DB 15; Length 410;
Best Local Similarity 99.3%; Pred. No. 1.8e-123;
Matches 405; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2551 ctcagaagctgtgtctcttgcctccacgcacgcacccgtatctccctccttgcgtg 2610
DB 410 CTCAGAAGCTGTGTCTCTCTGCCCCCAGCAGCAGCAGCTATCTGCCCTCTTGTGCTGT 351
QY 2611 agaagctgaagacacggtccccccaggagagcagctcagataggtgtgtgagcgtgc 2670
DB 350 AGAAGCTGAAGACGACGGTCCCCCAGGAGGAGCAGCTCAGGATAGGTGTGTGAGCTGTGC 291
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```
2y 2671 cgaggctgggtccacataaagcactagctctatagatgctcttaggaagctggtgctgg 2730
Db 290 CGAGGCTTGGGGTCCACATAGCACTAGTCTATAGATGCTCTTAGGACTGGTGGCTGG 231
-y 2731 cacagcggggcagagagctccacacgagcaagcagatgaactaatttcattca 2790
-b 230 CACAGCTGGGGCCAGAGAGCTCCACACGAGCAGCAGAGCACTAATTTCAITTC 171
2y 2791 aggcagtttttaagaagcttggaaacagacgcgccaccttctcttaataccagaaa 2850
Db 170 AGCAGTTTTTAAAGAAGTCATGGAACACAGCGCGGCACCTTCCCTTAATCCAGCAA 111
2y 2851 gtaattccctgcacacagacagacagcagatgaacagatcagtggtctaaagtccga 2910
-b 110 GTATTCCCTGCACACAGAGCAAGCAGAGTAACAGATCAGTGGGCTTAAGTGTCCGA 51
2y 2911 gacttaacgaaatagatttcagctgctcaataaagattgagttgcaa 2958
Db 50 GACTTAACGAAATAGTATTTCAGCTGCAATAAAGATTGAGTTGCAA 3

RESULT 65
LOCUS AW592223/c 461 bp mRNA EST 22-MAR-2000
DEFINITION bf41a01.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
IMAGE:2934408 3', mRNA sequence.
ACCESSION AW592223
VERSION AW592223.1 GI:7279399
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 461)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -40UP from Glibco
High quality sequence stop: 450.
Location/Qualifiers
1. .461
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2934408"
/clone_lib="Soares_NFL_T_GBC_S1"
/lab_host="DH10B"
/notes="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Equal amounts of plasmid DNA from three normalized
libraries (fetal lung NBH19W, testis NHT, and B-cell
NCI-CGAP-GCB1) were mixed, and ss circles were made in
vitro. Following HAP purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from pools of 5,000 clones made
from the same 3 libraries. The pools consisted of
I.M.A.G.E. clones 297480-302087, 682632-687239,
726408-728711, and 729096-731399. Subtraction by Bento
Soares and M. Fatima Bonaldo."
BASE COUNT 93 a 125 c 114 g 129 t
ORIGIN

Query Match 8.6%; Score 255; DB 92; Length 461;
Best Local Similarity 99.1%; Pred. No. 1.8e-123;
Matches 455; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

2500 agccacagcccaagaggtcagagccagctgaagatctgggagacccctgaactcagaag 2559
Db 461 AGCCACAGCCCAAGAGGTCAGAGCCAGTCGAAGATCTGGGAGACCCCTGAACCTCAGAGG 402
2y 2560 ctgtgtctcttctgccccacgcgcacccgctatctgcccctcttctgctgtagaagctga 2619
Db 401 CTGTGTCTCTTCTGCCCCACGCACGCCGTATCTGCCCTCTTGTGTGTAGAGCTGA 342
2y 2620 agagcagctccccccagggagcagctcagagataggtggatggatgagctgtgccagagcttg 2679
Db 341 AGAGCAGGCTCCCCAGAGGAGCAGCTCAGATAGGTGGTATGGAGCTGTGCCAGGCTTG 282
2y 2680 ggctccacataagcactagctctatagatgctctcttaggactgtgctggtgcacagcgc 2739
Db 281 GGTCCACATAGACACTAGTCTATAGATGCCCTCTTAGGACTGGTGGCTGGCAGAGCTGC 222
2y 2740 gggccagggaggtgccacacggaagcagatgaactaattcttcaaacggcagttt 2799
Db 221 GGCCAGGAGGCTGCCACACGGAAGCAAGCAGATGAACATAATTTCAATTTCAAGGCAGTTT 162
2y 2800 ttaaagaagctctggaaacagacgagggcgacaccttctcttaataccagcaagtgattccc 2859
Db 161 TTAAGAAGTCATGGAAACAGACGGCGGCACCTTCTCTTAATCCACAAATGATTCCC 102
2y 2860 tgcacaccagagacaagcagatgaacagatcagtggtggtctaaagtgtccgagacttaacg 2919
Db 101 TGCACACCAGAGACAGCAGAGTAACAGATCAGTGGGCTTAAGTGTCCGAGACTTAACG 42
2y 2920 aaatagatttcagctgcaataaagattgagttgcaa 2958
Db 41 AAAATAGTATTTCAGCTGCAATAAAGATTGAGTTGCAA 3

RESULT 66
LOCUS AA716607/c 479 bp mRNA EST 29-DEC-1997
DEFINITION 2968g07.s1 Soares_fetal_heart_NBHH19W Homo sapiens cDNA clone
IMAGE:398556 3', mRNA sequence.
ACCESSION AA716607
VERSION AA716607.1 GI:2728881
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 479)
AUTHORS Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisels, G., Jost, S.,
Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin,
J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B.,
White, Y., Wylie, T., Waterston, R. and Wilson, R.
WashU-NCI human EST Project
Unpublished (1997)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -40ml3 fwd. Et from Amersham
High quality sequence stop: 462.
Location/Qualifiers
1. .479
/organism="Homo sapiens"
/db_xref="GDB:1306379"
/db_xref="taxon:9606"
/clone="IMAGE:398556"
/clone_lib="Soares_fetal_heart_NBHH19W"
/sex="unknown"
/dev_stage="19 weeks"
/lab_host="DH10B (ampicillin resistant)"
/notes="Organ: heart; Vector: pT7T3D (Pharmacia) with a
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

Aufrey, C., Behar, G., Bois, F., Bouchier, C., da Silva, C., Devignes, M.D., Duprat, S., Houllatte, R., Jumeau, M.N., Lamy, B., Lorenzo, F., Mitchell, H., Mariage-Samson, R., Pietu, G., Pouillot, Y., Sebastiani-Kabaktchis, C. and Tessier, A.

TITLE

IMAGE: molecular integration of the analysis of the human genome and its expression

JOURNAL

C. R. Acad. Sci. III, Sci. Vie 318 (2), 263-272 (1995)

MEDLINE

95277534

COMMENT

Contact: Genzentrum Muenchen
Laboratorium fuer molekulare Biologie
Am Klopferplatz 18a, 8033 Martinsried, Germany
Email: obermaier@vms.biochem.mpg.de
single read.

FEATURES

Location/Qualifiers

1. 299

/organism="Homo sapiens"

/db_xref="GDB:D057384E"

/db_xref="taxon:9606"

/clone="HEI030"

/clone_lib="Stratagene cDNA library Human heart,

cat#936208"

/note="Vector: pBluescript SK(+); Human heart cDNA

library. Cloning vector pBluescript SK(+)"

BASE COUNT

96 a 77 c 70 g 56 t

ORIGIN

Query Match 8.4%; Score 248; DB 147; Length 299;
Best Local Similarity 99.7%; Pred. No. 9e-120;
Matches 298; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

451 caaagtgtactttctggacctccacactggaataatcacctcgaagcaatacaataat 510

|||||

2b 1 CAAAGTGTACTTTCTGGACCTCCACAACCTCGAATAATACCTCGAAGCAATCAAAATAT 60

|||||

451 ttcttggtccattgaaagaatagaactggtgtgctgcccccaactctgccccagataacg 570

|||||

2b 61 TTCTGGTCCATTGAAGGAATAGAACTGGCTGTGCGGCCCACTCTGCCCCAGAAATACT 120

|||||

451 aggatgaacatgacagtcttaccagatcccatcacacagtgaaacagagagggaaagc 630

|||||

2b 121 AGGATGAACCACTGACAGTTTACCAAGATCCCATACAGTGAACAGAGAGGGGAAGC 180

|||||

451 accaaccatggcagagtcacagaaggcctctcagcaggtcagtcagtcagcagcagcag 690

|||||

2b 181 ACCAACCATGGCAGAGTCCAGAAGGCTCTCAGCAGGCTCAGTCCAGAGCGATCTTCAG 240

|||||

451 actcagagtcgaataaataagacacaccttccacatggttttagccagagaagagg 749

|||||

2b 241 ACTCCGAGTCGAATGAAATGAGCCACACCTTCCACATGCTGTGTTAGCCAGAGAAGGG 299

|||||

RESULT

69

AI991599

LOCUS

AI991599/c

DEFINITION

ws18c04.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:2497542 3,

similar to SW:YATA_SCHPO Q10155 HYPOTHETICAL 90.6 KD PROTEIN

C1D4.10 IN CHROMOSOME 1.; mRNA sequence.

ACCESSION

AI991599

VERSION

AI991599.1

KEYWORDS

EST.

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 517)

AUTHORS

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

JOURNAL

COMMENT

Contact: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert_Strausberg@nih.gov

Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael

R. Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima

Bonaldo, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 625 Std Error: 0.00

Seq primer: -40Up from Gibco

High quality sequence stop: 426.

FEATURES

Location/Qualifiers

1. 517

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2497542"

/clone_lib="NCI_CGAP_GC6"

/tissue_type="pooled germ cell tumors"

/lab_host="DH10B"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified

polylinker; Plasmid DNA from the normalized library

NCI_CGAP_GC4 was prepared, and ss circles were made in

vitro. Following HAP purification, this DNA was used as

tracer in a subtractive hybridization reaction. The driver

was PCR-amplified cDNAs from a pool of 5,000 clones made

from the same library (cloneIDs 1257096-1258631,

1469064-1470983, and 1475592-1476743). Subtraction by

Bento Soares and M. Fatima Bonaldo."

BASE COUNT 118 a 139 c 123 g 136 t 1 others

ORIGIN

Query Match 8.4%; Score 247; DB 27; Length 517;

Best Local Similarity 100.0%; Pred. No. 3.1e-119;

Matches 247; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1846 tgcaccacatcagtatgattcctgcacaaatgccttcaggaagggtcagatctccagtc 1905

|||||

Db 447 TGCACCACATCAGTATGATTCCTGCCAAATGCTTCAGGAAGGGCTGAGATCTCCAGTC 388

|||||

Qy 1906 ctgcagtgaagaattgatcagttcgctgctgcgaacatgattggaagaagtttcaga 1965

|||||

Db 387 CTGCAGTGGAAAGATTGATCAGTTGCTGCTTCGGAACATGATTGGAAGAGTTTCAGA 328

|||||

Qy 1966 cctgtctggtgcgcactgcaagcatgcgtttgctgtgctgtgctgcacacctctgct 2025

|||||

Db 327 CTTGCTCTGCTGGGACATGTCAGCATGCTTGGCTGTGCGCTGTGCACACCTCTGGCT 268

|||||

Qy 2026 ggaagtggtctattcccggggacaccatgcctcgcagaggtctctgtccggatgggaaag 2085

|||||

Db 267 GGAAGTGTCTATTCCGGGACACCATGCTCGAGGCTCTGTCGGATGGGGAAG 208

|||||

Qy 2086 atgccac 2092

|||||

Db 207 ATGCCAC 201

|||||

RESULT

70

BE938229

LOCUS

BE938229

DEFINITION

CM4-TN0060-290800-565-B02 TN0060 Homo sapiens cDNA, mRNA sequence.

ACCESSION

BE938229

VERSION

BE938229.1

KEYWORDS

EST.

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 387)

AUTHORS Dias Neto, E., Garcia Correa, R., Verjowski-Almeida, S., Briones, M.R., Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordian, S., Costa, F.F., Goldman, G.H., Carvalho, A.F., Matsukuma, A., Baia, G.S., Simpson, D.H., Brunstein, A., deOliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.

TITLE Shotgun sequencing of the human transcriptome with ORF expressed sequence tags

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

MEDLINE 20202663

COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL (<http://www.ludwig.org.br/scripts/gethtml2.pl?tl=st2-CM4-TN0060-290>)
800-565-B02st3-2000-08-29st4-1)
Seq primer: puc 18 forward
High quality sequence stop: 385.

```

DEFINITION   oe91f04.s1 NCI_CGAP_Col2 Homo sapiens cDNA clone IMAGE:1419007, mRNA sequence.
ACCESSION    AA838624
VERSION      AA838624.1 GI:2913423
KEYWORDS     EST.
SOURCE       human.
ORGANISM     Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE    1 (bases 1 to 429)
              NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS      National Cancer Institute, Cancer Genome Anatomy Project (CGAP).
TITLE        Tumor Gene Index
JOURNAL
COMMENT      Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: Stratagene, Inc.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 1787 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 375.
              Location/Qualifiers
FEATURES             source
                    1..429
                     /organism="Homo sapiens"
                     /db_xref="taxon:9606"
                     /clone="IMAGE:1419007"
                     /clone_lib="NCI_CGAP_Col2"
                     /sex="mixed"
                     /tissue_type="colon tumor"
                     /lab_host="SOLR (kanamycin resistant)"
                     /note="Organ: colon; Vector: Bluescript SK-; Site_1: EcoRI ; Site_2: XhoI; Cloned unidirectionally. Primer: Oligo dt. Pooled colon tumors. 5' adaptor sequence: 5' GAATTCGGCCAGAG 3' adaptor sequence: 5' CTCGAGCTTTTTTTTTTTT3' Average insert size: 1.2 kb."
BASE COUNT      91 a 115 c 108 g 115 t

Query Match          8.3%; Score 245; DB 12; Length 429;
Best Local Similarity 99.5%; Pred. No. 3.6e-118;
Matches 415; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 2539 ggagaccctgaactcagaaggctgtgtgcttttcgccacgcagcgaccggtatctgcc 2598
DB 416 GGAGACCCTGAAC TCAGAAGGCTGTGTCTCTTGCCGCACGCCAGCCGCTATCTGCC 357
QY 2599 ctcttgtgttagaacctgaagcacggtctcccccaggagcgacctcagatatgggt 2658
DB 356 CTCTTGTGTTGAAGCTGAAGACGACGGTCCCCCAGAGGCGACTCAGATAGTGGT 297
QY 2659 atggagctgtgccagggttggtgggtcccaataagcactagtctatagatgccctttagg 2718
DB 296 ATGGAGCTGTGCCAGGGCTTGGGTCCCAACAATAAGCACTAGTCTATAGATGCCCTTAGG 237
QY 2719 actggtgcttgccacagccgcggccaggaggtctgccacacggaaagcagcatgaact-2778
DB 236 ACTGTTGCTT-GCACAGCCGGCGGCAGAGGCGTCCACACAGCAAGCAGATGAAC T 178
QY 2779 adtttcattcaaggcagctttttaagaagctcttggaaaacagcggcgccaccttcttc 2838
DB 177 AATTTCATTTCAAGGCAGTTTTTAAGAAAGTCATGGAACAGAGCGCGGCACCTTCTC 118
QY 2839 taatccagcaagtgtattccctgcacaccagacaagcagagtaacaggatcagtggtg 2898
DB 117 TAATCCAGCAAGTGATTCCCTGCACACCCAGACAGCAGAGTAACAGAGTACAGTGGGT 58

```



```

362 cctggaacacattctctgacacgaatgactggtctaatgttgggggcttaagtgaat 421
|||||
233 CTTGGACAAATATCTTGACAGCAATGCTGCTCTAATGTTGGGGCTTAAGTGAAT 174
|||||
422 gattcttactttaaagaaacccgggtctccaaagtgtgtactttctggacctccacaact 481
|||||
173 GATCTTACTTTAAAGGAACCGGGCTTCCAAAGTGTGTACTTCTGGACCTCCACAAC 114
|||||
482 ggaataatcctcgaagcaatcaaaaattttctgtgtccattgaagggaataagaactggc 541
|||||
113 GGAATAATACCTCGAAGCAATCAAAATATTTCTGTGCTCCATTGAAAGGAATAGAACTGGC 54
|||||
542 tctgcccgcacctctgcccagaataacagagatgaacacacacacatttacc 594
|||||
53 TGTGCGGCCCCACTGTCGCCAGAAACGAGGATGAAACCATGACAGTTTACC 1

RESULT 78
LOCUS AA522537/c 865 bp mRNA EST 20-AUG-1997
DEFINITION ni38e08.s1 NCI_CGAP_Lu1 Homo sapiens cDNA clone IMAGE:979142 3',
mRNA sequence.
ACCESSION AA522537
VERSION AA522537.1 GI:2263249
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 865)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R.
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: Stratagene, Inc., David B. Krizman,
Ph.D.
CDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/dbrrp/image/image.html
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 436.
FEATURES
source
1..865
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="NCI_CGAP_Lu1"
/tissue_type="lung tumor"
/lab_host="SOLR (kanamycin resistant)"
/notes="Organ: lung; Vector: Bluescript SK-; Site_1: EcoRI;
Site_2: XhoI; Cloned unidirectionally. Primer: Oligo dr.
Bulk lung tumor. 5' adaptor sequence: 5' GAATTCGGCAGAG 3'
3' adaptor sequence: 5' CTCGAGTTTTTTTTTTTTTTT 3'
Average insert size: 1.1 kb."
BASE COUNT 179 a 235 c 227 g 218 t 6 others
ORIGIN
Query Match 7.8%; Score 232; DB 8; Length 865;
Best Local Similarity 99.5%; Pred. No. 2.9e-111;
Matches 402; Conservative 0; Mismatches 1; Indels 1; Gaps 1;
2540 gagaccctgaactcagaagctgtgtgtcttctgcccacagcagcaccgctatctgccc 2599
|||||
403 GAGACCCTGAACCTCAGAAGGCTGTGTCTCTCTGCCACGACCGCATCTCTGCC 344
|||||

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QY 2600 tcttctgtagaagctgaagcacgcgtcccccaggaggcagctcagatagggtga 2659
|||||
Db 343 TCTTCTGCTAGAACCTGAAGACGCGGTCCCGGAGGAGGAGCTCAGATAGGTGGTA 284
|||||
QY 2660 tggagctgtgcccagagcttgggtctccacacataagcactagtctatagatgcctcttagga 2719
|||||
Db 283 TGGAGCTGTGCCAGGCTTGGGCTCCACATAGCACTAGTCTATAGATGCCCTTTAGGA 224
|||||
QY 2720 ctgggtgccttggcacagccgcgggcccaggggcgcacacggaagcaagcagatgaacta 2779
|||||
Db 223 CTGCTGCCCT-GCACACGCCGCGGCGCAGAGGCTGCACACGGAAGCAAGCAGATGAAC 165
|||||
QY 2780 attctattcaagcagcttttttaaaagaactcttgaaaacagcgcgcgcctcttctct 2839
|||||
Db 164 ATTTCATTTCAAGGCAAGTCTTTTAAAGAAGTCATGGAACAGACGGCGGCACCTTCTCT 105
|||||
QY 2840 aatccagcaaatgattccctgcacaccagagacacagcagagtaacagatcagtggtg 2899
|||||
Db 104 AATCCAGCAAGAGTGAATTCCTGCACACGAGACAGAGTAACAGGATCAGTGGGTC 45
|||||
QY 2900 taagtctcgcagacttaacgaaaataagttatttcagctgcaataa 2943
|||||
Db 44 TAAGTCTCCGAGACTTTAACGAAAATAGTATTTTCAGCTGCAATAA 1

RESULT 79
LOCUS AA928608/c 282 bp mRNA EST 07-JUL-1998
DEFINITION om75b03.s1 NCI_CGAP_GC4 Homo sapiens cDNA clone IMAGE:1552973 3',
mRNA sequence.
ACCESSION AA928608
VERSION AA928608.1 GI:3076899
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 282)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arraying by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/dbrrp/image/image.html
Insert Length: 846 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 255.
FEATURES
source
1..282
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1552973"
/clone_lib="NCI_CGAP_GC4"
/tissue_type="pooled germ cell tumors"
/lab_host="DH10B"
/notes="Vector: pTT73D-Pac (Pharmacia) with a modified
polylinker; 1st strand cDNA was prepared from 3 pooled
germ cell tumors, and was then primed with a Not I -
oligo(dT) primer. Double-stranded cDNA was ligated to Eco
RI adaptors (Pharmacia), digested with Not I and, cloned
into the Not I and Eco RI sites of the modified pT773
vector. Library is normalized. Library was constructed by
Bento Soares and M. Fatima Bonaldo."
BASE COUNT 63 a 68 c 66 g 84 t 1 others

```

ORIGIN

Query Match 7.7%; Score 228; DB 13; Length 282;
 Best Local Similarity 100.0%; Pred. No. 3.5e-109;
 Matches 228; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2y 2730 gcacagccgcccagagagctgcacacggaagcaagcagatgaactatcttc 2789
 |||||||
 2b 228 GCACAGCCGCGGCGAGAGCTGCCACACGGAAGCAAGCAGATGAATATTC 169
 |||||||
 2y 2790 aagcagcttttaagaagctttgaaacacagcgcgcaccccttccttaaccagcaa 2849
 |||||||
 2b 168 AAGCAGCTTTTAAAGAAGTCTTGGAAACACAGCGCGCACCTTTCTCTAATCAGCAA 109
 |||||||
 2y 2850 agtattccctgcacacagagacaagcagagtaacagatcagtggtctaaagtctcg 2909
 |||||||
 2b 108 AGTGATTCCTGCACACAGAGACAAGCAGAGTAACAGGATCAGTGGGTCTAAGTGTCCG 49
 |||||||
 2y 2910 agacttaacgaaatagatttcagctcgaataaagattgattgca 2957
 |||||||
 2b 48 AGACTTAACGAAATAGATTTCAGCTGCTCAATAAAGATTGAGTTGCA 1
 |||||||

RESULT 80

LOCUS T34024 282 bp mRNA EST 06-SEP-1995
 DEFINITION EST61387 Human White blood cells Homo sapiens cDNA 5' end similar
 to None, mRNA sequence.

ACCESSION T34024
 VERSION T34024.1 GI:616122

KEYWORDS

EST

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 282)

AUTHORS Adams, M.D., Kerlavage, A.R., Fleischmann, R.D., Fuldner, R.A., Bult

, C.J., Lee, N., Kirkness, E.F., Weinstock, K.G., Gocayne, J.D., White

, O., Sutton, G., Blake, J.A., Brandon, R.C., Chiu, M.-W., Clayton, R.A.,

Cline, R.T., Cotton, M.D., Earle-Hughes, J., Fine, L.D., FitzGerald

, L.M., FitzHugh, W.M., Fritchman, J.L., Geoghagen, N.S.M., Glodek, A.,

Ghosh, C.L., Hanna, M.C., Hedblom, E., Hinkle, J.P., Kelley, J.M.,

Klimek, K.M., Kelley, J.C., Liu, L.-I., Marmaros, S.M., Merrick, J.M.,

Moreno-Palances, R.F., McDonald, L.A., Nguyen, D.T., Pellegrino, S.M.,

Phillips, C.A., Ryder, S.E., Scott, J.L., Saudek, D.M., Shirley, R.,

Small, K.V., Spriggs, T.A., Utterback, T.R., Weidman, J.F., Li, Y.,

Bednarek, D.P., Cao, L., Cepeda, M.A., Coleman, T.A., Collins, E.-J.,

Dinke, D., Feng, P., Ferrie, A., Fischer, C., Hastings, G.A., He, W.-W.,

Hu, J.-S., Greene, J.M., Gruber, J., Hudson, P., Kim, A., Kozak, D.L.,

Kunsch, C., Ji, H., Li, H., Weissner, P.S., Olsen, H., Raymond, L., Wei

, Y.-F., Wing, J., Xu, C., Yu, G.-L., Ruben, S.M., Dillon, P.J., Fannon

, M.R., Rosen, C.A., Haseltine, W.A., Fields, C., Fraser, C.M. and

Venter, J.C.

Initial Assessment of Human Gene Diversity and Expression Patterns

Based Upon 83 Million Basepairs of cDNA Sequence

Nature 377, 3-174 (1995)

96026280

Other ESTs: THC15444

Contact: Venter, JC

The Institute for Genomic Research

932 Clopper Rd, Gaithersburg, MD 20878

Tel: 3018699056

Fax: 3018699423

Email: tdbinfo@tdb.tigr.org

For clone availability, additional sequence and expression

information related to this EST, please contact the TIGR Database

(tdbinfo@tdb.tigr.org)

Seq primer: M13 Reverse.

Location/Qualifiers

1. 282

/organism="Homo sapiens"

/db_xref="ATCC (inhost):104694"

FEATURES

source

/db_xref="taxon:9606"
 /clone_lib="Human White blood cells"
 /tissue_type="white blood cells"
 /note="organ: blood"

BASE COUNT 62 a 83 c 62 t 2 others
 ORIGIN

Query Match 7.7%; Score 228; DB 145; Length 282;
 Best Local Similarity 100.0%; Pred. No. 3.5e-109;
 Matches 228; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1672 ctgtgtttgtctccacacctgcacgcagatcacccacacggcttgccaatctctgtgc 1731
 |||||||
 Db 26 CTGTGTTGTCTCCACCTGTCAGCAGATCACACACGGGCTTGCCAGATATCTTGTCTGC 85
 |||||||
 Qy 1732 agagagaacgcgcttggcattcttgggaagcgcgcttcaccccttgcgtggtgccc 1791
 |||||||
 Db 86 AGAGAGAACGGCCTTGGCATCTTTGGCAAAAGCCCTTCACCCCTTGTGTGGTGGTGC 145
 |||||||
 Qy 1792 ccaaccagctcaagcctggctccagcagatcaccaacacagtcgacagaggtctgcacc 1851
 |||||||
 Db 146 CCAACCACTCAAGCCTGGCTCCAGCAGTACCAACAGTCGACGAGGTCCTGCACC 205
 |||||||
 Qy 1852 acatcagtatgattctcccaaatgcttcagggaagggcgtgagatct 1899
 |||||||
 Db 206 ACATCAGTATGATTCCTGCCAAATGCCCTTCAGGAAGGGCTGAGATCT 253
 |||||||

RESULT 81

AI357786/c

LOCUS AI357786

DEFINITION QU98d07.x1 NCI-CCAP_Gas4 Homo sapiens cDNA clone IMAGE:1980109 3',

mRNA sequence.

AI357786

VERSION AI357786.1

KEYWORDS EST

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 433)

REFERENCE NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.

AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

CONTACT Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

CDNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CCAP clone distribution, information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 2467 Std Error: 0.00

Seq primer: -400P from Gibco

High quality sequence stop: 397.

Location/Qualifiers

1. 433

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1980109"

/clone_lib="NCI-CCAP Gas4"

/tissue_type="poorly differentiated adenocarcinoma with

signet ring cell features"

/lab_host="DH10B"

/note="organ: stomach; Vector: pCMV-SPORT6; Site:1: SalI;

Site:2: NotI; Cloned unidirectionally. Primer: Oligo dT.

Average insert size 1.69 kb. Life Technologies catalog #:

11549-011"

AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Oligo-dT track not found, Not I site shown in beginning of sequence
is likely internal to the message. cDNA Library Preparation: M.B.
Soares Lab clone distribution: NCI-CGAP clone distribution
information can be found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Seq primer: M13 Forward
POLYA-NO.

FEATURES
source

Location/Qualifiers
1. .228
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2730521"
/clone_lib="NCI-CGAP_Sub6"
/lab_host="DH10B (Life Technologies)"
/note="vector: pT73B-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; NCI-CGAP_Sub6
is a subtracted library derived from BW, which consists of
a mixture of four normalized libraries: NCI-CGAP_Brn50,
NCI-CGAP_Lu13, NCI-CGAP_Ov18, GBC1. The NCI-CGAP_Sub6
library had 7 million recombinants. A single-stranded DNA
preparation of BW was used as a tracer in a subtractive
hybridization with a driver comprising: the IMAGE pool
(NCI-CGAP_Kid3 pool 1 LLAM 3334-3337, 3682-3683,
3798-3803 (IMAGE Clones 1323376-1323911,
1456008-1456775, 1500552-1502855); NCI-CGAP_Kid5 pool 1
LLAM 3338-3342, 3722-3725, 3776-3778 (IMAGE Clones
1323912-1325831, 1471368-1472903, 1492104-1493255);
NCI-CGAP_Lu5 pool 1 LLAM 3575-3582, 3851-3854 (IMAGE
Clones 1414920-1417991, 1520904-1522439); NCI-CGAP_GC4
pool 1 LLAM 3164-3167, 3716-3720, 3733-3735 (IMAGE
Clones 1257096-1258631, 1469064-1470983, 1475592-1476743
); NCI-CGAP_P22 pool 1 LLAM 2457-2459, 2758-2759,
3062-3068 (IMAGE Clones 985608-986759, 1101192-1101959,
1217928-1220615); NCI-CGAP_Co10 pool 1 LLAM 2644-2653,
2871-2872 (IMAGE Clones 1057416-1061255, 1144584-1145351
). (50% of the driver population), plus a pool of 3,840
arrayed clones from NCI-CGAP_Sub1 (IMAGE Clones
2708618-2710535) and NCI-CGAP_Sub2 (IMAGE Clones
2710536-2712455) (20% of the driver population), plus a
pool of 11,136 clones from NCI-CGAP_Sub3 (IMAGE Clones
2712456-2723591) (30% of the driver population).
Subtraction was performed as previously described [Bonaldo
, Lennon & Soares (1996): Normalization and Subtraction:
Two Approaches To Facilitate Gene Discovery. Genome
Research 6, 791-806.
TAG_LIB=NCI-CGAP-Lu13
TAG_TISSUE=lung
TAG_SEQ=GCCGG"
BASE COUNT 41 a 72 c 78 g 37 t
ORIGIN

Query Match 7.4%; Score 218; DB 88; Length 228;
Best Local Similarity 100.0%; Pred. No. 5.9e-104;
Matches 218; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
1 CGGCGCGCAAGGACCCGCTGCGGACCTGCGCAGAGAGAGCGGACCGCTCGGG 60
141 CGGCGCGCAAGGACCCGCTGCGGACCTGCGCAGAGAGAGCGGACCGCTCGGG 200
1 CGGCG 60
201 TGCTCG 260
61 TGCTCG 120
261 GCGCGCGCGCTACGCTTCTCCGAGTTCACCGGATCTCTTCACTGTGGAAGGC 320

Db 121 GGCGCGCGCTCTAGCTTCTCTCGAGTTCACCGGTATCTCTTCACTGTGGAGAGC 180
Qy 321 gttcagagactcgcaggagacacaaagttaaagggttc 358
|||||
Db 181 GTTCAGAGACTCATGCGAGGACACAAAGTTAAAGGTTC 218
RESULT 86
N36229 448 bp mRNA EST 16-JAN-1996
LOCUS Y330C04.s1 Soares melanocyte 2NDHM Homo sapiens cDNA clone
DEFINITION IMAGE:272742 3' similar to SW:YK59.YEAST P36159 HYPOTHETICAL 96.8
KD PROTEIN IN SIS2-MTD1 INTERGENIC REGION. [1] ; mRNA sequence.
N36229
ACCESSION N36229.1 GI:1157371
VERSION N36229.1
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 448)
AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman,
M., Hulman, M., Kucaba, T., Le, M., Lennon, G., Mairra, M., Parsons, J.,
Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaskis, E., Waterston,
R., Williamson, A., Wohldmann, P. and Wilson, R.
TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Wilton RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
High quality sequence stops: 365
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (infoimage.llnl.gov) for further information.
Possible reversed clone: similarity on wrong strand
Possible reversed clone: polyT not found
Seq primer: ml3 -40 forward
High quality sequence stop: 365.
FEATURES
Location/Qualifiers
1. .448
/organism="Homo sapiens"
/db_xref="GDB:3882384"
/db_xref="taxon:9606"
/clone="IMAGE:272742"
/clone_lib="Soares melanocyte 2NDHM"
/sex="Male"
/tissue_type="melanocyte"
/lab_host="DH10B (ampicillin resistant)"
/note="vector: pT73B (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was primed with a Not I - oligo(dT) primer [5,
TGTTACCAATCTGAAGTGGGAGCGCGGCGGAGTTTGTGTTT 3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pT73 vector
(Pharmacia). Library constructed by Bento Soares and
M.Fatima Bonaldo. RNA from normal foreskin melanocytes
(FS374) was kindly provided by Dr. Anthony P. Albino."

BASE COUNT 110 a 113 c 127 g 94 t 4 others
ORIGIN
Query Match 7.4%; Score 218; DB 142; Length 448;
Best Local Similarity 100.0%; Pred. No. 7.2e-104;
Matches 218; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 252 cgggactcggcgccgcgctcagctcttcctccagttcaaccggtatctcttcaactgt 311
Db 109 CGGAGACTGGCGCGCGCGCTCTAGCTTCTCTCCGAGTTCACCGGTATCTCTCACTGT 168


```

2b 159 TGGAAACACACGGCGACCTTCTCTAATCCAGCAAGAGTATTCCTCCGACACACAGAG 100
2y 2872 acaagcagagtaacaggatcagtggtctaaagtgtccgagacttaacgaaatagtattt 2931
2b 99 ACAAGCAGAGTAAACAGGATCAGTGGGTCTTAAGTCCGAGACTTAACGAAAAATAGTATT 40
2y 2932 cagctgcaataaagattgagttgcaa 2958
2b 39 CAGCTGCAATAAAGATTGAGTTTGCAA 13

RESULT 91
LOCUS W37591 422 bp mRNA EST 10-OCT-1996
DEFINITION zc10f03.r1 Soares parathyroid_tumor_NbHPA Homo sapiens cDNA clone
ACCESSION W37591
VERSION W37591.1 GI:1319196
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 422)
AUTHORS Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
R., Williamson,A., Wohlmann,P. and Wilson,R.
The WashU-Merck EST Project
Unpublished (1995)
CONTACT: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 678 Std Error: 0.00
Seq primer: mob.REGA+ET.
FEATURES
Location/Qualifiers
1..422
/organism="Homo sapiens"
/db_xref="GDB:1259575"
/db_xref="taxon:9606"
/clone="IMAGE:321917"
/tissue_type="parathyroid_tumor_NbHPA"
/dev_stage="adult"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: parathyroid gland; Vector: pTT73D (Pharmacia
) with a modified polylinker; Site:1: Not I; Site:2: Eco
RI; 1st strand cDNA was primed with a Not I - oligo(dT)
primer
[5'-TGTTACCAATCTGAAGTGGGCGCGCCACCAATTTTTTTTTTTTTTTT
TTTT-3'], double-stranded cDNA was size selected, ligated
to Eco RI adaptors (Pharmacia), digested with Not I and
cloned into the Not I and Eco RI sites of a modified pTT73
vector (Pharmacia). Library went through one round of
normalization to a Cot = 5. Library constructed by Bento
Soares and M.Fatima Bonaldo. RNA from sporadic parathyroid
adenomas was kindly provided by Dr. Stephen Marx, National
Institute of Diabetes and Digestive and Kidney Diseases,
NIH."
BASE COUNT 100 a 115 c 102 g 104 t 1 others
ORIGIN
Query Match 7.0%; Score 206; DB 146; Length:422;
Best Local Similarity 100.0%; Pred. No. 1.6e-97;
Matches 206; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
2y 841 tccagttgggacagctgccatgcctcccatcattgctgctgcaaggggaaagca 900

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Db 271 TCCCAAGTGGACAGCTGCCATCGCTCCCATCATTTGCTGCTCAAGGACGGAAAGCA 212
Qy 901 tcaactgaaggaagagagattttggtggaagagctgtgtactctccagatcctggtg 960
Db 211 TCACATCATGAAGGAAGAGAGATTTGGCTGAAGAGCTGTGTACTCTCCAGATCCTGGTG 152
Qy 961 ctgctttgtggtgtagaagtccagatgaagcttcaaccatctgtgagaatg 1020
Db 151 CTGCTTTTGTGGTGTAGTAATGTCAGATGAAGACTTATTCACCCATCTGTGAGATG 92
Qy 1021 ccaccttcagaggtaccaggaaag 1046
Db 91 CCACCTTCAGAGGTACCAAGGAAG 66

RESULT 92
LOCUS R51138 472 bp mRNA EST 18-MAY-1995
DEFINITION YG71c08.r1 Soares infant brain 1NIB Homo sapiens cDNA clone
IMAGE:38752.5' similar to SP:YK59_YEAST P36159 HYPOTHETICAL 96.8 KD
PROTEIN IN SIS2-MTD1 INTERGENIC ;, mRNA sequence.
ACCESSION R51138
VERSION R51138.1 GI:813040
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 472)
AUTHORS Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
R., Williamson,A., Wohlmann,P. and Wilson,R.
The WashU-Merck EST Project
Unpublished (1995)
CONTACT: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
High quality sequence stops: 327
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: M13RP1
High quality sequence stop: 327.
FEATURES
Location/Qualifiers
1..472
/organism="Homo sapiens"
/db_xref="GDB:411293"
/db_xref="taxon:9606"
/clone="IMAGE:38752"
/clone_lib="Soares infant brain 1NIB"
/dev_stage="73 days post natal"
/note="Organ: whole brain; Vector: Lqmid BA; Site:1: Not
I; Site:2: Hind III; 1st strand cDNA was primed with a Not
I - oligo(dT) primer [5'
AACGGAAGATTCGCGCCGCGCCAGGAATTTTTTTTTTTTTTTT 3'];
double-stranded cDNA was ligated to Hind III adaptors
(Pharmacia), digested with Not I and directionally cloned
into the Not I and Hind III sites of the Lqmid BA vector.
Library went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
BASE COUNT 105 a 131 c 125 g 102 t 9 others
ORIGIN
Query Match 6.7%; Score 199; DB 144; Length 472;
Best Local Similarity 100.0%; Pred. No. 8.4e-94;

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Matches 199; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1430 gcaggagtagagagtagcgcagagcagccagccagccagcagagagagagagtagta 1489
    |||||
    43 GCAGGAGTACAGGAGGAGTGCAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGT 102
QY 1490 ccagagaatacatcttctctggaacagaggtgtgcatccagcagagattcgaatgtcag 1549
    |||||
    25 CCCAGAAATCATCTTCTCTGGAACAGGTTGCCATCCCGATGAAGATTCCGAATGTCAG 162
QY 1550 tgcacacttgtcaacataagcccccacacagctctctgctactgactgtgtgagggcac 1609
    |||||
    25 TGCACACTTGTCAACATAGCCCGACACGCTCTGCTACTGAGCTGTGGTGAGGGCAC 222
QY 1610 atttggcagctgtgcgct 1628
    |||||
    23 ATTTGGGCGAGCTGTGCGGT 241

RESULT 93
LOCUS AI141263/c 416 bp mRNA EST 05-OCT-1998
DEFINITION qa46h05.s1 Soares_NhMPu_S1 Homo sapiens CDNA clone IMAGE:1689849
3', mRNA sequence.
ACCESSION AI141263
VERSION AI141263.1 GI:3648720
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 416)
AUTHORS C.J., Lee, N.H., Kirkness, E.F., Weinstein, K.G., Gocayne, J.D., White
TITLE NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
NATIONAL Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
UNPUBLISHED (1997)
CONTACT: Robert Strausberg, Ph.D.
TEL: (301) 496-1550
EMAIL: Robert.Strausberg@nih.gov
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 711 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 233.
FEATURES
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            /db_xref="taxon:9606"
            /clone="IMAGE:1689849"
            /clone_lib="Soares_NhMPu_S1"
            /tissue_type="Pooled human melanocyte, fetal heart, and
            pregnant uterus"
            /lab_host="DH10B"
            /note="Organ: mixed (see below); Vector: pT7A3D-Pac
            (Pharmacia) with a modified polylinker; Site 1: Not I;
            Site 2: Eco RI. Equal amounts of plasmid DNA from three
            normalized libraries (melanocyte 2NBHM, pregnant uterus
            NBHPU, and fetal heart NBHH19W) were mixed, and ss circles
            were made in vitro. Following HAP purification, this DNA
            was used as tracer in a subtractive hybridization
            reaction. The driver was PCR-amplified cDNAs from pools of
            5,000 clones made from the same 3 libraries. The pools
            consisted of i.M.A.G.E. clones 260232-265223,
            340488-345479, and 484488-489479."
            87..a 108 c 99 g 122 t
            /ase_count
            /origin

Query Match 6.7%; Score 198; DB 16; Length 416;
Best Local Similarity 99.0%; Pred. No. 2.8e-93;
Matches 398; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
2557 aggtgtgtgtctctgccccagcagcagcccgatctctgcccctctgtgtgtagaagc 2616

```

```

Db 416 AGCTGTGTCTTCTGCCCCAGCAGCAGCCCGATCTGCCCCCTCTGCTGGTAGAAGC 357
QY 2617 tgaagagcagcgggtccccccagggagcagcagcagcagcagcagcagcagcagcagc 2676
    |||||
    356 TGAAGAGCAGCGTCCCGCAGGAGGAGCCTCAGATAGGTGGTATGGAGCTGTGCCAGGC 297
QY 2677 ttggggtcccaataagaagcactagctatagatgcctctcttaggactggttcctggcagcagc 2736
    |||||
    296 TTGGGTCCCCACATAGCAGCTAGTCTATATAGTGCCTCTTAGGACTGGTGCCTGGCACAGC 237
QY 2737 cgcggccagcaggggtgcccacgcaggaagcaagcagatgaactaatttcatttcagagcag 2796
    |||||
    236 TGGGGCCAGGAGGCTGCCACAGCGAGGAGCAAGCAGATGAATTAATTTCAATTTCAAGGGCAG 177
QY 2797 ttttaagaagctcttggaaacagcagcggcgccctctctcttaataccagcagcagcagc 2856
    |||||
    176 TTTTAAAGAGTCATGTGAAGAACAGACAGCGCGGCGACCTTCTCTAATCCAGCAAAATGATT 117
QY 2857 cctgcacacagcagcagcagcagcagcagcagcagcagcagcagcagcagcagcagcagc 2916
    |||||
    116 CCTGTCCACACAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGC 57
QY 2917 acgaaaatagttttcagctgcaataaagatgagttgtcaa 2958
    |||||
    56 ACGAAATAGTATTCAGCTGCCAATAAAGATTGAGTTGCCAA 15

RESULT 94
LOCUS AA346268 394 bp mRNA EST 21-APR-1997
DEFINITION EST52407 Greater omentum IV Homo sapiens CDNA 5' end, mRNA
sequence.
ACCESSION AA346268
VERSION AA346268.1 GI:1998525
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 394)
AUTHORS Adams, M.D., Kerlavage, A.R., Fleischmann, R.D., Fuldner, R.A., Bult
, C.J., Lee, N.H., Kirkness, E.F., Weinstein, K.G., Gocayne, J.D., White
, O., Sutton, G., Blake, J.A., Brandon, R.C., Man-Wai, C., Clayton, R.A.,
Cline, T.R., Cotton, M.D., Earle-Hughes, J., Fine, L.D., Fitzgerald
, L.M., Fitzhugh, W.M., Fritchman, J.L., Geoghegan, N.S., Glodek, A.,
Gnehm, C.L., Hanna, M.C., Hedblom, E., Hinkle, P.S., Jr., Kelley, J.M.,
Kelley, J.C., Liu, L.-I., Marmaros, S.M., Merrick, J.M.,
Moreno-Palancas, R.F., McDonald, L.A., Nguyen, D.T., Pelligrino, S.M.,
Phillips, C.A., Ryder, S.E., Scott, J.L., Saudek, D.M., Shirley, R.,
Small, K.V., Spriggs, T.A., Utterback, T.R., Weidman, J.F., Li, Y.,
Bednarek, D.P., Cao, L., Cepeda, M.A., Coleman, T.A., Collins, E.J.,
Dimke, D., Feng, D.-F., Ferrie, A., Fischer, C., Hastings, G.A., He, W.W.,
Hu, J.S., Greene, J.M., Gruber, J., Hudson, P., Kim, A.K., Kozak, D.L.,
Kunsch, C., Hung, J., Li, H., Meissner, P.S., Olsen, H., Raymond, L.,
Wei, Y.F., Wing, J., Xu, C., Yu, G.L., Ruben, S.M., Dillion, P.J., Fannon
, M.R., Rosen, C.A., Haseltine, W.A., Fields, C., Fraser, C.M., and
Venter, J.C.
TITLE Initial assessment of human gene diversity and expression patterns
based upon 83 million nucleotides of cDNA sequence
NATURE 377 (6547 Suppl), 3-174 (1995)
MEDLINE 96026280
COMMENT Other_ESTs: HCL175624
Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA
Tel: 3018699056
Fax: 3018699423
Email: arkerlavage@tigr.org
For clone availability, additional sequence and expression
information related to this EST, please check the TIGR Human Gene
Index (http://www.tigr.org/tdb/hgi/hgi.html)

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FEATURES
  source
    Seq primer: (-21)ML3_universal.
    Location/Qualifiers
      1..290
        /organism="Homo sapiens"
        /db_xref="taxon:9606"
        /clone="c-23e06"
        /clone_lib="normalized infant brain cdna"
        /sex="Female"
        /tissue_type="total brain"
        /dev_stage="3 months old"
        /note="Organ: brain; Vector: lafmid BA; Site_1: HindIII;
        Site_2: NotI; sex:Female; dev_stage=3 months old;
        isolate=muscular atrophy patient; tissue_type=total brain
        ; total mRNA was oligo (dT) primed and directionally
        cloned 5' -> 3' into the HindIII -> NotI sites of the
        lafmid BA vector. Clone library from B.Soaes, Psychiatry
        Dept. Columbia University, USA. Normalization_method:
        Bento Soares, P.N.A.S in press"
      BASE COUNT      65 a      72 c      80 g      70 t      3 others
      ORIGIN
        Query Match      6.2%; Score 182; DB 147; Length 290;
        Best Local Similarity 100.0%; Pred. No. 8.3e-85;
        Matches 182; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

      1026 ttccagggtaccagaagaaagcagatgccccctggccttgggtgttcacatggcccca 1085
      109 TTTTCCAGAGGTACCAAGAAAGCAGATGCCCCCGTGGCTTGGTGGTTTCATATGGCCCCA 168
      1086 gcatctgtgctgtggacagcagatcaccagcagtgatggagaggttggcctgacacc 1145
      169 GCATCTGTGCTGTGGACAGCAGTACCAGCAGTGGATGGAGAGGTTTGGCCCTGACACC 228
      1146 cagcacttggctcctgaatgagaactgtgctcagttcacacacccctgcagccacaagatt 1205
      229 CAGCACTTGGTCTGTAATGAGAACTGTGCTCAGTTTCAACACCTTCGCGAGCCACAAGATT 288

      1206 ca 1207
      289 CA 290

      RESULT 97
      LOCUS H03318 424 bp mRNA EST 20-JUN-1995
      DEFINITION YJ47el0.s1 Soares placenta NB2HP Homo sapiens cDNA clone
      IMAGE:151914 3', mRNA sequence.
      ACCESSION H03318
      VERSION H03318.1 GI:866251
      KEYWORDS EST.
      SOURCE human.
      ORGANISM Homo sapiens
      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
      Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
      1 (bases 1 to 424)
      Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman
      M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,
      Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevisan, E., Waterston
      R., Williamson, A., Wohldmann, P. and Wilson, R.
      The WashU-Merck EST Project
      Unpublished (1995)
      Contact: Wilson RK
      Washington University School of Medicine
      4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
      Tel: 314 286 1800
      Fax: 314 286 1810
      Email: est@wustl.edu
      Insert Size: 1152
      High quality sequence stops: 315
      Source: IMAGE Consortium, LLNL
      This clone is available royalty-free through LLNL; contact the
      IMAGE Consortium (info@image.llnl.gov) for further information.

FEATURES
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    Insert Length: 1152 Std Error: 0.00
    Seq primer: Promega -2mln13
    High quality sequence stop: 315.
    Location/Qualifiers
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        /db_xref="GDB:564054"
        /db_xref="taxon:9606"
        /clone="IMAGE:151914"
        /clone_lib="Soares placenta NB2HP"
        /sex="Female"
        /dev_stage="placenta obtained at birth (full term)"
        /lab_host="DH10B (ampicillin resistant)"
        /note="Organ: placenta; Vector: pTZ19 (Pharmacia) with a
        modified polylinker; Site_1: NotI; Site_2: EcoRI; 1st
        strand cDNA was primed with a NotI - oligo(dT) primer [5',
        AACTGGAGAAATTCGCGCGCAGGAAATTTTTTTTTTTT 3'],
        double-stranded cDNA was ligated to EcoRI adaptors
        (Pharmacia), digested with NotI and cloned into the NotI
        and EcoRI sites of the modified pTZ19 vector. Library
        went through one round of normalization. Library
        constructed by Bento Soares and M.Fatima Bonaldo.
      BASE COUNT      89 a      105 c      116 g      106 t      8 others
      ORIGIN
        Query Match      6.2%; Score 182; DB 141; Length 424;
        Best Local Similarity 100.0%; Pred. No. 8.5e-85;
        Matches 182; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

      2762 aagcaagcagatgaactattcattcaggcagtttttaagaaagctcttgaaacaga 2821
      195 AGCAACGACATGAATTAATTCATTTCAGGCGAGTTTAAAGAGACTCTTGGAACAGA 136
      2822 cggcgccaccttctctcttaataccagcaaatgattccctgcacaccagagacagcag 2881
      135 CGCGCGCACCTTCTCTTAATCCAGCAAGTATTCCCTGCACACCAGACAGCAGAG 76
      2882 taacagagatcagtggtcttaagtgtccgagacctaagaaatagtttcagctgcaat 2941
      75 TAACAGGATCAGTGGGTCTAAGTGTCCGAGACTTAACGAAATAGTATTTCAGCTGCAAT 16
      2942 aa 2943
      15 AA 14

      RESULT 98
      LOCUS AI033342/c 482 bp mRNA EST 28-AUG-1998
      DEFINITION OX02d12.s1 Soares fetal_liver_spleen_inFLS_S1 Homo sapiens CDNA
      clone IMAGE:165159 3', mRNA sequence.
      ACCESSION AI033342
      VERSION AI033342.1 GI:3254295
      KEYWORDS EST.
      SOURCE human.
      ORGANISM Homo sapiens
      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
      Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
      1 (bases 1 to 482)
      NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
      Tumor Gene Index
      Unpublished (1997)
      Contact: Robert Strausberg, Ph.D.
      Tel: (301) 496-1550
      Email: Robert.Strausberg@nih.gov
      This clone is available royalty-free through LLNL; contact the
      IMAGE Consortium (info@image.llnl.gov) for further information.
      Insert Length: 749 Std Error: 0.00
      Seq primer: -40ml3 fwd. ET from Amersham
      High quality sequence stop: 455.
      Location/Qualifiers

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1. 482
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:165159"
/clone_lib="Soares_fetal_liver_spleen_lNFLS_S1"
/sex="male"
/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: Liver and Spleen; Vector: pT73D (Pharmacia)
with a modified polylinker; Site_1: Pac I; Site_2: Eco RI;
This is a subtracted version of the original Soares fetal
liver spleen lNFLS library. 1st strand cDNA was primed
with a Pac I - oligo(dT) primer [5',
AAGCGAAGATTAATTAAGATCTTTTTTTTTTTTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
BASE COUNT      94 a 130 c 122 g 136 t
ORIGIN
Query Match      6.1%; Score 180; DB 15; Length 482;
Best Local Similarity 100.0%; Pred. No. 9.9e-84;
Matches 180; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2502 ccacagggcgaaggtcagagccagtcgaagatctggagaccctgaactcagaaggtc 2561
      |||
JB 456 CCACAGGCCAAGAGGTCAGAGCCAGTGAAGATCTGGGAGACCCCTGAAGAGGCT 397

QY 2562 ggtgtctcttcccccacagcagcaccctatctgcctcttctggttagagctgaag 2621
      |||
JB 396 GGTGTCTCTTCCGCCACACGACGCCCTATCTGCCCTCTTGTGGTAGAGCTGAAG 337

QY 2622 agcacggtcccccagagcagtcagtcagtagtggtgtagctgcccagggcttgg 2681
      |||
JB 336 AGCACGGTCCCCAGGAGGACGCTCAGGATAGGTGTTATGGAGCTGTGCCGAGGCTTGG 277

RESULT 99
LOCUS      BE619874      577 bp      mRNA      EST      24-AUG-2000
DEFINITION 6014731301 NIH_MGC_68 Homo sapiens cDNA clone IMAGE:3876223 3',
      mRNA sequence.
ACCESSION  BE619874
VERSION     BE619874.1 GI:9890812
KEYWORDS   EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 577)
AUTHORS     NIH-MGC http://www.ncbi.nlm.nih.gov/MGC/.
TITLE       National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL     Unpublished (1999)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Tel: (301) 496-1550
            Email: Robert.Strausberg@nih.gov
            Tissue Procurement: DCTD/DPG/Gazdar
            CDNA Library Preparation: Life Technologies, Inc.
            CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone Distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLC623 row: k column: 08
            High quality sequence stop: 577.
            Location/Qualifiers
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                /db_xref="taxon:9606"
                /clone="IMAGE:3876223"

FEATURES
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1. 482
/organism="Homo sapiens"
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/clone="IMAGE:165159"
/clone_lib="Soares_fetal_liver_spleen_lNFLS_S1"
/sex="male"
/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: Liver and Spleen; Vector: pT73D (Pharmacia)
with a modified polylinker; Site_1: Pac I; Site_2: Eco RI;
This is a subtracted version of the original Soares fetal
liver spleen lNFLS library. 1st strand cDNA was primed
with a Pac I - oligo(dT) primer [5',
AAGCGAAGATTAATTAAGATCTTTTTTTTTTTTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
BASE COUNT      94 a 130 c 122 g 136 t
ORIGIN
Query Match      6.1%; Score 180; DB 15; Length 482;
Best Local Similarity 100.0%; Pred. No. 9.9e-84;
Matches 180; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2502 ccacagggcgaaggtcagagccagtcgaagatctggagaccctgaactcagaaggtc 2561
      |||
JB 456 CCACAGGCCAAGAGGTCAGAGCCAGTGAAGATCTGGGAGACCCCTGAAGAGGCT 397

QY 2562 ggtgtctcttcccccacagcagcaccctatctgcctcttctggttagagctgaag 2621
      |||
JB 396 GGTGTCTCTTCCGCCACACGACGCCCTATCTGCCCTCTTGTGGTAGAGCTGAAG 337

QY 2622 agcacggtcccccagagcagtcagtcagtagtggtgtagctgcccagggcttgg 2681
      |||
JB 336 AGCACGGTCCCCAGGAGGACGCTCAGGATAGGTGTTATGGAGCTGTGCCGAGGCTTGG 277

RESULT 100
LOCUS      T72963      376 bp      mRNA      EST      02-MAR-1995
DEFINITION yc5b06.r1 StrataGene liver (#937224) Homo sapiens cDNA clone
      IMAGE:85523 5', mRNA sequence.
ACCESSION  T72963
VERSION     T72963.1 GI:689638
KEYWORDS   EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 376)
AUTHORS     Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiapelli, B.,
            Chissoe, S., Dietrich, N., Dubucque, T., Favello, A., Gish, W., Hawkins
            M., Hultman, M., Kucaba, T., Lacy, M., Le, N., Mardis, E., Moore
            B., Morris, M., Parsons, J., Prange, C., Rifkin, L., Rohlfing, T.,
            Schellenberg, K., Soares, M.B., Tan, F., Thierri-Meg, J., Trevisakis, E.,
            Underwood, K., Wohlmann, P., Waterston, R., Wilson, R. and Maizra, M.
            Generation and analysis of 280,000 human expressed sequence tags
            Genome Res. 6 (9), 807-828 (1996)
            9704478
            Contact: Wilson RK
            Washington University School of Medicine
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: est@watson.wustl.edu
            Insert Size: 954
            High quality sequence stop: 335.
            Location/Qualifiers
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                /db_xref="taxon:9606"
                /clone="IMAGE:85523"
                /clone_lib="StrataGene liver (#937224)"
                /sex="male"
                /dev_stage="49 years old"
                /lab_host="SOLR cells (kanamycin resistant)"

FEATURES
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/db_xref="GDB:502580"
/db_xref="taxon:9606"
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/clone_lib="StrataGene liver (#937224)"
/sex="male"
/dev_stage="49 years old"
/lab_host="SOLR cells (kanamycin resistant)"

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/note="Organ: liver; Vector: pBluescript SK; Site_1: EcoRI
; Site_2: XhoI; Cloned unidirectionally. Primer: Oligo
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size: 1.1 kb; Uni-ZAP XR Vector; -5' adaptor sequence: 5'
GAATTCGGCAGCAG 3' -3' adaptor sequence: 5'
CTCGAGTTTTTTTTTTTTT 3'"

BASE COUNT 82 a 94 c 121 g 63 t 16 others

ORIGIN

Query Match 6.0%; Score 178; DB 145; Length 376;
Best Local Similarity 100.0%; Pred. No. 1.1e-82;
Matches 178; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
2208 attatgtaaacacattcagccagcgctatgccaaggtcccccctcttcagcccaacttc 2267
|||||
37 ATTATGCTGAACCACTTCAGCCAGCGCTATGCCAAGGTCCCCCTCTTCAGCCCAACTTC 96
2268 agcgagaaagtggagttgccttgaccacatgaaggtctgttgagagactttccaaca 2327
|||||
97 AGCGAGAAAGTGGAGTTGCTTTGACCACATGAAGTCTGCTTTGGAGACTTTCACACA 156
2328 atgcccagctgattcccccaactgaagccctgtttgctgagacatcgaggagatgg 2385
|||||
157 ATGCCAAGCTGATTCCCCCACTGAAGCCCTGTTTCTGGCGACATCGAGGAGATGG 214

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